VOLUME 1.

THEY KAR BOOK FOR

COLORISTS & DYERS

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RARE BOOKCASE

THE YEAR-BOOK

FOR

COLORISTS AND DYERS

Presenting a Review of the Year's Advances in the Bleaching, Dyeing, Printing, and Finishing of Textiles

RY

HARWOOD HUNTINGTON

VOLUME I

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2417

PREFACE.

So far as I am aware, there are no portable works in the English language to which a color-chemist can refer and find the information which he requires the oftenest.

The need of a concise recapitulation of advances made annually in the special field worked in by dyers and colorists has long been felt. New books and literature come along in such wheelbarrow-loads, and the number of periodicals has increased to such a large extent, that it is difficult for anyone who does not make a specialty of the subject to keep abreast with the times.

The object of this year-book is to meet this demand with accuracy and brevity. Such work, thoroughly and conscientiously done, cannot do otherwise than produce a book of value to those working in color-chemistry.

It is not meant to replace the larger books, but simply to serve as a *vade meeum*. The compiler will be abundantly repaid if the industry is advanced in some measure by this publication.

It is hoped that kind friends will call attention to any mistakes or any omissions, so that these may be remedied in the issue of next year.

HARWOOD HUNTINGTON.

NEW YORK CITY, October, 1898.





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THURST HITCH

PART I.—TABLES.

COMPARISON OF METRIC SYSTEM WITH THE UNITED STATES METHOD OF WEIGHTS AND MEASURES.

(Arranged in Alphabetical Order.)

Are (100 square meters) = 119.6 square yards.

Bushel = 2150.42 cubic inches, 35.24 liters.

Centare (1 square meter) = 1,550 square inches.

Centigram (1/100 gram) = 0.1543 grain.

Centiliter (1/100 liter) = 2.71 fluid drams, 0.338 fluid ounce. Centimeter (1/100 meter) = 0.3937 inch.

1 Cubic centimeter = 16.23 minims (Apothecaries).

10 " centimeters = 2.71 fluid drams (Apothecaries).

30 " - " = 1.01 " ounces "
100 " = 3.38 " " "
473 " = 16.00 " " "
500 " = 16.90 " " "
1.000 " = 33.81 " " "

Decigram (1/10 gram) = 1.5432 grains.

Decimeter (1/10 meter) = 3.937 inches.

Deciliter (1/10 liter) = 0.845 gill.

Dekagram (10 grams) = 0.3527 ounce.

Dekaliter (10 liters) = 9.08 quarts (dry), 2.6418 gallons.

Dekameter (10 meters) = 393.7 inches.

Dram = 1.772 grams.

Dram (Apothecaries or Troy) = 3.9 grams.

Foot = 0.3048 meter, or 30.48 centimeters.

Gallon = 4,543 liters.

Gill = 0.118295 liter, or 142 cubic centimeters.

Grain (Troy) = 0.064804 gram.

Grain = 0.0648 gram.

Gram = 15.432 grains.

Hectare (10,000 square meters) = 2.471 acres.

Hectogram = 3.5274 ounces.

Hectoliter (100 liters) = 2.838 bushels, or 26.418 gallons.

Hectometer (100 meters) = 328 feet 1 inch.

Hundredweight (112 pounds Avoirdupois) = 50.8 kilograms.

Inch = 0.0254 meter.

Inch = 2.54 centimeters.

Inch = 25.40 millimeters.

Kilogram = 2.2046 pounds, or 35.274 ounces.

Kiloliter (1,000 liters) = 1.308 cubic yards, or 264.18 gallons.

Kilometer (1,000 meters) = 0.62137 mile (3,280 feet 10 inches).

Liter = 1.0567 quarts, 0.264 gallon (liquid), or 0 908 quart (dry).

Meter = 39.3704 inches, or 3.28087 feet.

Mile = 1.609 kilometers.

Mile = 5,280 feet, or 1609.3 meters.

Millier or tonneau = 2,204.6 pounds.

Milligram = 0.0154 grain.

Millimeter (1/1,000 meter) = 0.0394 inch.

Myriagram = 22.046 pounds.

Myriameter (10,000 meters) = 6.2137 miles.

Ounce (Avoirdupois) = 28.350 grams.

Ounce (fluid) = 28.3966 cubic centimeters.

Ounce (Troy or Apothecaries) = 31.104 grams.

Ounce (Avoirdupois) = 28.35 grams.

Peck = 9.08 liters.

Pint (liquid) = 0.47318 liter (liquid), or 0.568 (dry).

Pound (Avoirdupois) = 453.603 grams.

Pound (English) = 0.453 kilogram.

Pound (Troy) = 373.25 grams.

Quart = 1.1352 liters.

Quart (liquid) = 0.94636 liter.

Quintal = 220.46 pounds.

Scruple (Troy) = 1.296008 grams.

Ton = 20 hundredweight = 2,240 pounds (Avoirdupois) 1016.070 kilograms.

Yard = 0.9144 meter.

APPROXIMATE EQUIVALENTS.

Centimeter, about 1/3 inch.

Cubic centimeter, about a thimbleful.

Dime, about 21/2 grams.

Dollar, silver, weighs about 25 grams. Five-cent nickel, about 5 grams. Kilogram, about 2 pounds. Kilometer, about $\frac{1}{2}$ mile. Liter, about a quart. Meter, about a yard.

LONG MEASURE.

```
12
    inches (in.) = 1 foot (ft.)
3
    feet
               =1 yard (yd.)
5½ yards
               =1 rod (rd.)
    rods
40
               =1 furlong (fur.)
    furlongs
              =1 mile (m.)
8
3
    miles
               =1 league
```

SURFACE OR SQUARE MEASURE.

```
144 square inches = 1 square foot.
9 square feet = 1 square yard
301/4 square yards = 1 square rod or perch
160 square rods = 1 acre
```

Measure 209 feet on each side, and you have a square acre approximately.

CUBIC MEASURE.

```
1,728 cubic inches (cu. in.) = 1 cubic foot (cu. ft.)
27 cubic feet = 1 cubic yard (cu. yd.)
40 cubic feet of hewn timber | = 1 ton or load (t.)
50 cubic feet = 1 cord foot (cd. ft.)
8 cord feet, or | = 1 cord of wood (cd.)
```

LIQUID OR WINE MEASURE.

```
4 gills (gi.) = 1 pint (pt.)
2 pints = 1 quart (qt.)
4 quarts = 1 gallon (gal.)
```

DRY MEASURE.

2 pints (pt.) = 1 quart (qt.) 8 quarts = 1 peck (pk.) 4 pecks = 1 bushel (bu.)

APOTHECARIES' WEIGHT.

20 grains = 1 scruple 3 scruples = 1 dram 8 drams = 1 ounce 12 ounces = 1 pound

AVOIRDUPOIS, OR COMMERCIAL WEIGHT.

27.34375 grains = 1 dram

16 drams = 1 ounce = 4371/2 grains

16 ounces = 1 pound = 256 drams = 7,000 grains.

28 pounds = 1 quarter = 448 ounces

4 quarters = 1 hundredweight = 112 pounds

20 hundredweights = 1 ton = 80 quarters = 2,240 pounds.

The standard of the avoir pound, which is the one in common commercial use, is the weight of 27.7015 cubic inches of pure distilled water, at its maximum density at about 39° F., in latitude of London, at the level of the sea, barometer at 30". But this involves an error of about 1 part in 1,362, for the 1 pound of water == 27.68122 cubic inches.

A troy pound = 0.82286 avoir pound. An avoir pound = 1,21528 troy pound, or apothecaries.

A troy ounce = 1.09714 avoir ounce. An avoir ounce = 0.911458 troy ounce or apothecaries.

Where we use the word "ton" we always mean 2,240 pounds, because that is its meaning in United States law.

MISCELLANEOUS WEIGHTS.

Barrel of flour = 196 pounds.
Barrel of salt = 280 pounds.
Bale of cotton (in America) = 400 pounds.

Bale of cotton (in Egypt) = 90 pounds. Bag of Sea Island cotton = 300 pounds.

Cable = 120 fathoms.

Cask of lime = 240 pounds.

Fathom = 6 feet.

Hand = 4 inches.

Hogshead = 63 gallons.

Keg (nails) = 100 pounds.

Pace = 3.3 feet.

Palm = 3 inches.

Pipe = 2 hogsheads.

Stone = 14 pounds.

Tun = 2 pipes.

Cubic foot of water weighs 62.4 pounds.

Cubic foot of water is 7.48 gallons.

Gallon of water weighs 81/2 pounds.

Gallon of water is 1,728 eubic inches.

In England, wool is sold by the sack, or boll, of 22 stones, which, at 14 pounds the stone, is 308 pounds.

A pack of wool is 17 stones and 2 pounds, which is rated as a pack load for a horse. It is 240 pounds.

A tod of wool is 2 stones of 14 pounds.

A wey of wool is 61/4 tods. Two weys, a sack.

A clove of wool is half a stone.

TABLE OF MULTIPLES.

Centimeters \times 0.3937 = inches.

Centimeters \times 0.0328 = feet.

Centimeters, cubic, \times 0.0338 = apothecaries' fluid ounces

Diameter of a circle × 3.1416 = circumference.

Gallons \times 3.785 = liters.

Gallons \times 0.833565 = imperial gallons.

Gallons \times 8.33505 = pounds of water.

Gallons, imperial, \times 1.199666 = U. S. gallons.

Gallons, imperial, \times 10 = pounds of water.

Gallons, imperial, \times 4.54102 = liters.

Grains \times 0.0648 = grams.

Inches \times 0.0254 = meters.

Inches \times 25.4 = millimeters.

Miles × 1.609 = kilometers.

Ounces, Troy, × 1.097 = ounces of avoirdupois.

Ounces, avoirdupois, × 0.9115 = ounces Troy.

Pounds, avoirdupois, × 0.4536 - kilograms.

Pounds, avoirdupois, × 0.8228571 = pounds Troy.

Pounds, Troy, × 0.37286 = kilograms.

Pounds, Troy, × 1.21527 = pounds avoirdupois.

Radius of a circle × 6.283185 = circumference.

Square of the radius × 3.1416 = area.

Square of the diameter of a circle × 0.7854 = area.

Square of the circumference of a circle × 0.07958 = area.

APPROXIMATE METHOD FOR ROUND VATS.

Rule.—Multiply the depth in feet by the diameter in feet, and that product by $1\frac{1}{2}$, and you have the number of barrels of $31\frac{1}{2}$ gallons each.

ANOTHER RULE FOR THE MEASUREMENT OF CYLINDRICAL CISTERNS.

Take the length, width, and depth in feet; multiply these together, and the product by 1,865; cut off four figures on the right, and the result will be the contents in barrels.

THERMOMETRY.

Fahr.	Centigr.	Fahr.	Centigr.	Fahr.	Ceat'gr.
-10	-23.33	2	-16.67	14	-10.00
9	-22.78	3	-16.11	15	- 9.44
- 8	-22.22	4	-15.56	16	- 8.89
— 7	-21.67	5	15.00	17	- 8.33
— 6	21.11	6	-14.44	18	— 7.7 3
— 5	-20.56	7	-13.89	19	— 7.22
4	-20.00	8	-13.33	20	- 6.67
— 3	-19.44	9	-12.78	21	— 6.11
— 2	-18.80	10	-12.22	22	- 5.56
- 1	-18.33	11	-11.67	23	- 5.00
0	-17.78	12	-11.11	24	- 4.44
1	-17.22	13	-10.56	25	- 3.89

TABLES.

Fahr.	Centigr.	Fahr.	Centigr.	Fahr.		Centigr.
26	- 3.33	66	18.89	106		41.12
27	- 2.78	67	19.44	107	t	41.67
28	- 2.22	68	20.00	108		42.23
29	- 1.67	69	20.56	109		42.78
30	- 1.11	70	21.11	110		43.34
31	- 0.56	71	21.67	111		43.89
32	- 0.00	72	22.22	112	;	44.45
33	0.56	. 73	22.78	113		45.00
34	1.11	74	23.33	114		45.56
35	1.67	75	23.89	115		46.22
36	2.22	76	24.44	116	:	46.67
37	2.78	77	25.00	117		47.23
38	3.33	78	25.56	118		47.78
39	3.89	79	26.11	119		48.34
40	4.44	80	26.67	120	*	48.89
41	5.00	81	27.22	121		49.45
42	5.56	82	27.78	122		50.00
43	6.11	83	28.33	123	•	50.56
44	6.67	84	28.89	124	1	51.12
45	7.22	85	29.44	125		51.67
46	7.78	86	30.00	126		52.23
47	8.33	87	30.56	127		52.78
48	8.89	88	31.11	128		53.34
49	9.44	89	31.67	129		53.89
50	10.00	90	32.22	130		54.45
51	10.56	91	32.78	131		55.00
52	11.11	92	33.33	132		55.56
53	11.67	93	33.89	133		56.12
54	12.22	94	34.44	134		56.67
55	12.78	95	35.00	135	•	57.23
56	13.33	96	35.56	136		57.78
57	13.89	97	36.11	137		58.34
58	14.44	98	36.67	138		58.89
59	15.00	99	37.22	139		59.45
60	15.56	100	37.78	140		60.00
61	16.11	101	38.34	141		60.56
62	16.67	102	38.89	142		61.12
63	17.22	103	39.45	143		61.67
64	17.78	104	40.00	144		62.23
65	18.33	105	40.56	145		62.78

				P	
Fahr.	Centigr.	Fahr.	Centigr.	Fahr.	Centigr.
146	63.34	187	86.12	227	108.33
147	63.89	188	86.67	228	108.89
148	64.45	189	87.23	229	109.44
149	65.00	190	87.78	230	110.00
150	65.56	191	88.34	231	110.55
151	66.12	192	.88.89	232	111.11
152	66.67	193	89.45	233	111.67,
153	67.23	194	90.00	234	112.22
154	67.78	195	90.56	235	112.78
155	68.34	196	81.12	236	113.33
156	68.89	197	91.67	237	113.89
157	69.45	198	92.23	238	114.44
158	70.00	199	92.78	239	115.00
159	70.56	200	93.34	240	115.55
160	71.12	201	93.89	241	116.11
161	71.67	202	94.44	242	116.67
162	72.23	203	95.00	243	117.22
163	72.78	204	95.55	244	117.78
164	73.34	205	96.11	245	118.33
165	73.89	206	96.67	246	118.89
166	74.45	207	97.22	247	119.44
167	75.00	208	97.78	248	120.00
168	75.56	209	98.33	249	120.55
169	76.12	210	98.89	250	121.11
170	76.67	211	99.44	251	121.67
171	77.23	212	100.00	252	122.22
172	77.78	213	100.55	253	122.78
173	78.34	214	101.11	254	123.33
174	78.89	215	101.67	255	123.89
175	79.45	216	102.22	256	124.44
176	80.00	217	102.78	257	125.00
177	80.56	218	103.33	258	125.55
178	81.12	219	103.89	259	126.11
179	\$1.67	220	104.44	260	126.67
180	82.23	221	105.00	261	127.22
181	82.78	222	105.55	262	127.78
182	83.34	223	106.11	263	128.33
183	83.89	224	106.67	264	128.89
184	84.45	225	107.22	265	129.44
185	85.00	226	107.78	266	130.00
186	85.56				

Another Method to Transform Fahrenheit to Centigrade, or vice versa, is the formula:

$$\frac{\mathrm{C} \, \times \, 9}{5} \, + \, 32$$
 equals Fahrenheit.

$$F-32 \times \frac{5}{9}$$
 equals Centigrade.

Another Way to Convert Fahrenheit to Centigrade Degrees, subtract 32 and divide by 2; then add to this 1/10 of itself, and, if further accuracy is desired, 1/100 more.

Comparison Between the Scales of Fahrenheit, Reaumur and the Centigrade.

(Zero Fahrenheit corresponds with minus 17.78 Centigrade and minus 14.22 Réaumur.)

Cent.	Fahr.	Rmr.	Cent.	Fahr.	Rmr.
100	212	80	79	174.2	63.2
99	210.2	79.2	78	172.4	62.4
98	208.4	78.4	77	170.6	61.6
97	206.6	77.6	76	168.8	60.8
96	204.8	76.8	75	167	60
95	203	76	74	165.2	59.2
94	201.2	75.2	73	163.4	58.4
93	199.4	74.4	. 72	161.6	57.6
92	197.6	73.6	71	159.8	56.8
91	195.8	72.8	70	158	56
90	194	72	69	156.2	55.2
89	192.2	71.2	68	154.4	54.4
88	190.4	70.4	67	152.6	53.6
87	188.6	69.6	66	150.8	52.8
86	186.8	68.8	65	149	52
85	185	68	64	147.2	51.2
84	183.2	67.2	63	145.4	50.4
83	181.4	66.4	62	143.6	49.6
82	179.6	65.6	61	141.8	48.8
81	177.8	64.8	60	140	48
80	176	64	59	138.2	47.2

Cent.	Fahr.	Rmr.	Cent.	Fahr.	Rmr.
58	136.4	46.4	18	64.4	14.4
57	134.6	45.6	17	62.6 .	13.6
56	132.8	44.8	16	60.8	12.8
55	131	44	15	59	12
54	129.2	43.2	14	57.2	11.2
53	127.4	42.4	13	55.4	10.4
52	125.6	41.6	12	53.6	9.6
51	123.8	40.8	11	51.8	8.8
50	122	40	10	50	8
49	120.2	39.2	9	48.2	7.2
48	118.4	38.4	s	46.4	6.4
47	116.6	37.6	7	44.6	5.6
46	114.8	36.8	6	42.8	4.8
45	113	36	5	41	4
44	111.2	35.2	4	39.2	3.2
43	109.4	34.4	3	37.4	2.4
42	107.6	33.6	2	35.6	1.6
41	105.8	32.8	1	33.8	0.8
40	104	32	Zero	32	Zero
39	102.2	31.2	1	30.2	0.8
38	100.4	30.4	2	28.4	1.6
37	98.6	29.6	3	26.6	2.4
36	96.8	28.8	4	24.8	3.2
35	95	28	5	23	4
34	93.2	27.2	6	21.2	4.8
33	91.4	26.4	7	19.4	5.6
32	89.6	25.6	8	17.6	6.4
31	87.8	24.8	9	15.8	7.2
30	86	24	10	14	8
29	84.2	23.2	11	12.2	8.8
28	82.4	22.4	12	10.4	9.6
27	80.6	21.6	13	8.6	10.4
26	78.8	20.8	14	6.8	11.2
25	77	20	15	5	12
24	75.2	19.2	16	3.2	12.8
23	73.4	18.4	17	1.4	13.6
22	71.6	17.6	18	-	14.4
21	69.8	16.8	19	2.2	15.2
20	68	16	20	4	16
19	66.2	15.2	21	5.8	16.8

Cent.	Fahr.	Rmr.	Cent.	Fahr.	Rmr.
22	7.6	17.6	36	32.8	28.8
23	9.4	18.4	37	34.6	29.6
24	11.2	19.2	38	36.4	30.4
25	13	20	39	38.2	31.2
26	14.8	20.8	40	40	32
27	16.6	21.6	41	41.8	32.8
28	18.4	22.4	42.	43.6	33.6
29	20.2	23.2	43	45.4	34.4
30	22	24	44	47.2	35.2
31	23.8	24.8	45	49	36
32	25.6	25.6	46	50.8	36.8
33	27.4	26.4	47	52.6	37.6
34	29.2	27.2	48	54.4	38.4
35	31	28	49	56.2	39.2

SULPHURIC ACID.

The Manufacturing Chemists' Association have adopted the following figures as representing the per cent. of H_2SO_4 in an oil of vitriol. Since colorists rarely or never need to look up an acid below 80 per cent. oil of vitriol, the table only goes to that degree.

Beaumé.	Sp. Gr.	H ₂ SO ₄ .	Oil Vitriol.
66	1.835	93.50	100
65	1.814	88.82	95
64	1.793	86.02	92
63	1.767	83.21	89
62	1.747	81.34	87
61	1.727	79.47	85
60	1.705	77.60	83
59	1.683	75.73	81

AREOMETRY OR HYDROMETRY.

COMPARISON BETWEEN THE SPECIFIC GRAVITY OF BEAUMÉ AND TWADDLE.

Tw.	В.	Sp. Gr.	Tw.	В.	Sp. Gr.	Tw.	В.	Sp. Gr.	Tw.	В.	Sp. Gi
0	0	1,000	44	26.0	1,220	88	44 1	1.440	132	57 4	1.660
1	0.7	1.005	45	26.4	1.225	89	44.4	1.445	133	57.7	1.665
2	1.4	1.010	46	26 9	1.230	90	44.8	1.450	134	57.9	1.670
2	2.1	1.015	47	27.4	1.235	91	45.1	1.455	135	58.2	1.675
4	2.7	1.020	48	27.9	1.240	92	45.4	1.460	136	58.4	1.680
5	3.4	1.025	49	28.4	1.245	93	45.8	1.465	137	58.7	1.685
6	4.1	1.030	50	28.8	1.250	94	46.1	1.470 1.475	138	58.9	1.690
6	4.7	1 035	51	29.3	1.255 1.260 1.265	95	46.4	1.475	139	59.2	1.695
8	5.4	1.040	52	29.7	1.260	96	46.8	1.480	140	59.5	1.695 1.700
8 9	6.0	1.045	53	30.2	1.265	97	47.1	1.480 1.485	141	59.7	1 -05
10	6.7	1.050	54	30.6	1.270	98	47.4	1.490	142	60.0	1.710
11	7.4	1.055	55	31,1	1.275	99	47.8	1.490 1.495	143	60 2	1 715
12	8.0	1.060	56.	31.5	1.270 1.275 1.280 1.285	100	48.1	1.500	144	60.4	1.720
13	8.7	1.065	57	32.0	1.285	101	48.4	1.505	145	60,6	1,725
14	9.4	1.070	58	32.4	1.290	102	48.7	1.510	146	60.9	1.730
15	10.0	1.075	59	32.4 32.8	1.290 1.295	103	49.0	1.515	147	61.1	1.735
16	10,6	1.080	60	33.3	1.300	104	$\frac{49.4}{49.7}$	1.510 1.515 1.520 1.525	148	61.4	1.710 1.715 1.725 1.725 1.730 1.735 1.740
17	11.2	1.085	61	33,7	1.305	105	49.7	1.525	149	61.6	
18	11.9	1,090	62	34.2	1.310	106	50.0	1.550	150	61.8	1.750
18 19	12.4	1.095	63	34.6	1.315	107	50.3	1.535	1151	62.1	1,755
20 -	13,0	1.100	64	35.0	1.320	108	50.6	1.540	152	62.3	1.760
21 22	13.6	1.105 1.110 1.115 1.120	65	35,4	1.325 1.330 1.335	109 110	50.9 51.2 51.5 51.8	1.545	153	62.5	1, 765
55	14.2	1.110	66	35.8	1.330	110	51.2	1.550	154	62.8	1.770
23	14.9	1.115	67 68	36.2	1.335	111	51.5	1.555	155	63.0	1.775
24	15.4	1.120	68	36.6	1.340	112	51.8	1.560	156	63.2	1.775 1.780
25 26	16.0	1.125 1.130	69 70	37.0	1.345	113	52.1	1.565	157	C3.5	1.785
26	16.5	1.130	70	37.4	1.345 1.350	114	52.1 52.4 52.7 53.0	1.565 1.570 1.575 1.580 1.585	158	63.7	1.785 1.790
27 28	17.1 17.7	1.135 1.140	71	37.8	1.355	115	52.7	1.575	159	64.0	1.795
28	17.7	1.140	72	50.~	1.360	116	53.0	1.580	160	64.2	1.800
29	18.3	1.145	73	38.6	1.365	117	55.3	1.585	161	64.4	1.805 1.810 1.815
30	18.8	1.150 1.155	74 75	39.0	1.370	118	53.6	1.590	162	64.6	1.810
31	19.3	1.155	75	39.4	1.375	119	53.9	1.595	163	64.8	1.815
32 33	19.8	1.160 1.165 1.170 1.175	76 77	39.8	1.380	120 121	54.1	1,600	164	65.0	1.820 1.825 1.830 1.835
33	20.3	1.165	77	40.1	1.385	121	54.4	1.605	165	65.2	1.825
34	20.9	1.170	78	40.5	1.390	122	54.7	1.610	166	65.5	1.830
35	21.4	1.175	79	40.8	1.395	123	55.0	1.615	167	65.7	1.835
36 37	22.0	1.180	80	41.2 41.6	1.400	124 125	55.2	1.620 1.625 1.630	168	65.9	1.840
37	22.5	1.185	81	41.6	1.405	125	55.5	1.625	169	66.1	1.840 1.845 1.850 1.855
38 39	23.0	1.190	82 83	42.0	1.410	126	55.8	1.630	170	66.3	1.850
39	23.5	1.195	53	42.3	1.415	127	56.0	1,655	171	66.5	1.855
40	24.0	1.200	84	42.7	$1.420 \\ 1.425$	128	56.3	1.640 1.645	172	66.7	1.800
41	24.5	1.205	85	43.1	1.425	129	56.6	1.645	173	67.0	1.865
42	25.0	1.210	86	43.4	1.430	130	56.9	1.650			
43	25.5	1.215	87	43.8	1,435	131	57.1	1.655			

If the sample is too viscous to determine the density directly, a weighed portion of it can be first diluted with a weighed quantity of water, or a weighed portion can be dissolved and diluted to a known volume with water. In the first instance the per cent. of total solids is to be calculated by the following formula:

Per cent. of solids in the undiluted material $=\frac{WS}{W}$

S = per cent. of solids in the diluted material.

W = weight of the diluted material.

w = weight of the sample taken for dilution.

If the dilution was made to a definite volume, the following formula is to be used:

Per cent. of solids in the undiluted material $=\frac{\text{VDS}}{\text{w}}$.

V = volume of the diluted solution.

D = specific gravity of the diluted solution.

S = per cent. of solids in the diluted solution.

w = weight of the sample taken for dilution.

To reduce a liquid to a desired specific gravity:

Let v equal the volume of the first liquid.

Let B " Beaumé of the first liquid.

Let b " Beaumé of the second or reducing liquid.

Let a " " Beaumé required.

Let z " " volume of the reducing liquid.

$$z = \frac{av - vB}{b - a}.$$

TO CHANGE FROM TWADDLE TO SPECIFIC GRAVITY,

OR

TO CHANGE FROM SPECIFIC GRAVITY TO TWADDLE.

In order to change degrees Twaddle into specific gravity, multiply by 5, add 1,000, and divide by 1,000. *Example:* Change 168° Twaddle into specific gravity.

To change specific gravity into degrees Twaddle, multiply by 1,000, subtract 1,000, and divide by 5. Example: Change 1.84 specific gravity into degrees Twaddle.

The degrees of Twaddle's hydrometer are easily turned into specific gravity numbers-a quality which makes it preferable to any other hydrometer in use. The rule is to multiply the indicated degree by 5, and add 1,000 to the product; for example, 9 degrees Twaddle equals specific gravity 1,045; 25 degrees Twaddle equals specific gravity 1,125; 100 degrees Twaddle equals specific gravity 1,500; and so on. To bring specific gravity numbers to degrees of Twaddle, subtract 1,000, and divide the remainder by 5; for example, specific gravity 1,100 equals 20 degrees Twaddle.

WATER ANALYSIS.

Analyses of water are sometimes reported as parts per million, and sometimes grains per imperial gallon, and sometimes grains per United States gallon. The following table gives all three values:

Parts per Million.	Grains per Imperial Gallon.	Grains per United States Gallon.	Parts per Million.	Grains per Imperial Gallon.	Grains per United States Gallon.
1	0.0700	0.0583	13	0.9100	0.7581
2	0.1400	0.1166	14	0.9800	0.8165
3	0.2100	0.1749	15	1.0500	0.8747
4	0.2800	0.2332	16	1.1200	0.9330
5	0.3500	0.2915	17	1.1900	0.9914
6	0.4200	0.3499	18	1.2600	1.0497
7 .	0.4900 -	0.4082	19	1.3300	1.1080
8	0.5600	0.4665	20	1.4000	1.1663
9	0.6300	0.5248	21	1.4700	1.2246
10	0.7000	0.5831.	22	1.5400	1.2829
11	0.7700	0.6414	23	1.6100	1.3413
12	0.8400	0.6998	24	1.6800	1.3996

Parts per Million.	Grains per Imperial Gallon,	Grains per United States Gallon,	Parts per Million.	Grains per Imperial Gallon.	Grains per United States Gallon.
25	1.7500	1.4579	63	4.4100	3.6740
26	1.8200	1.5162	64	4.4800	3.7323
27	1.8900	1.5745	65	4.5500	3.7909
28	1.9600	1.6329	66	4.6200	3.8489
29	2.0300	1.6912	67	4.6900	3.9073
30	2.1000	1.7495	68	4.7600	3.9656
31	2.1700	1.8078	69	4.8300	4.0239
32	2.2400	1.8661	70	4.9000	4.0822
33	2.3100	1.9244	71	4.9700 .	4.1405
34	2.3800	1.9828	72	5.0400	4.1988
35	2.4500	2.0411	73	5.1100	4.2575
36	2.5200	2.0994	74	5.1800	4.3155
37	2.5900	2.1577	75	5.2500	4.3738
38	2.6600	2.2160	. 76	5.3200	4.4321
39	2.7300	2.2745	77	5.3900	4.4904
40	2.8000	2.3327	78	5.4600	4.5488
41	2.8700	2.3910	79	5.5300	4.6071
42	2.9400	2.4493	_80	5.6000	4.6654
43	3.0100	2.5076	81	5.6700	4.7237
44	3.0800	2.5659	82	5.7400	4.7820
45	3.1500	2.6243	83	5.8100	4.8403
46	3.2200	2.6826	84	5.8800	4.8987
47	3.2900	2.7409	85	5.9500	4.9570
48	3.3600	2.7992	86	6.0200	5.0154
49	3.4300	2.8575	87	6.0900	5.0736
50	3.5000	2.9129	88	6.1600	5.1319
51	3.5700	2.0742	89	6.2300	5.1903
52	3.6400	3.0325	90	6.3000	5.2486
53	3.7100	3.0908	91	6.3700	5.3069
54	3.7800	3.1491	92	6.4400	5.3652
55	3.8500	3.2074	93	6.5100	5.4235
56	3.9200	3.2658	94	6.5800	5.4818
57	3.9900	3.3241	95	6.6500	5.5402
58	4.0600	3.3824	96	6.7200	5.5985
59	4.1300	3.4407	97	6.7900	5.6568
60	4.2000	3.4990	98	6.8600	5.7151
61	4.2700	3.5573	99	6.9300	5.7734
62	4.3400	3.6157	100	7.0000	5.8318

TABLE OF PRINCIPAL ATOMIC MASSES.

Name.	Symbol.	Atomic Mass.	Name.	Symbol.	Atomic Mass.
Aluminum.	-	27.00	Lithium	•	7.02
Antimony	Sb	120.00	Magnesiun		24.30
Arsenic		75.00	Manganese	Mn	55.00
Barium	Ba	137.00	Mercury	Hg	200.00
Bismuth	Bi	208.00	Molybdenu	ımMo	96.00
Boron	B	11.00	Nickel	Ni	58.70
Bromin	\dots Br	79.95	Nitrogen	N	14.03
Cadmium	Cd	112.00	Oxygen	0	16.00
Calcium	Ca	40.00	Phosphoru	ıs P	31.00
Carbon	C	12.00	Platinum.	Pt	195.00
Chlorin	Cl	35.45	Potassium	K	39.11
Chromium.	Cr	52.10	Silicon	Si	28.40
Cobalt	Co	59.00	Silver	Ag	107.92
Copper	Cu	63.60	Sodium	Na	23.05
Fluorin	F	19.00	Strontium	Sr	87.60
Gold	Au	197.00	Sulfur	S	32.06
Hydrogen.	Н	1.00	Tin	Sn	119.00
Iodin	I	126.85	Tungsten	W	184.00
Iron	\dots Fe	56.00	Vanadium	V	51.40
Lead	Pb	206.95	Zine	Zn	65.41

PRONUNCIATION AND ORTHOGRAPHY OF CHEMICAL TERMS.

ADOPTED BY THE AMERICAN ASSOCIATION FOR THE ADVANCEMENT OF SCIENCE IN 1891.

alū minum.	chro mium.	nī trogen.
a'ntimony.	cō'balt.	ŏ'xygen.
a rsenic.	co'pper.	phos'phorus.
bā'rium.	flū orīn.	plă tinum.
bi'smuth (biz).	hy drogen.	potă ssium.
bō ron.	ī odĭn.	sī licon.
bro'mĭn.	iron.	silver.
că dinium.	lead.	sō'dium.
că lcium.	magnē'sium (zhium).	strö'ntium (shium).
carbon.	ma'nganese (eze).	sŭ lfur.
cē rium.	me rcury.	tin.
cē'sium.	môlý bdenum.	vănā dium.
chlö'rĭn.	nĭ ckel.	zinc.

TERMINATIONS IN -id (FORMERLY -ide).

The final e is dropped in every case and the syllable pronounced id as chlō'rĭd, ĭ'odĭd, h \bar{y} 'drĭd, ŏ'xĭd, h \bar{y} drŏ'xĭd, sŭ'lfĭd, ā'mĭd, ā'milīd.

TERMINATIONS IN -ane, -ene, -ine, AND -one.

The vowel of these syllables is invariably long, as me'thane, e'thane, na'phthalene, a'nthracene, pro'pine, qui'none, a'cetone, ke'tone,

MISCELLANEOUS WORDS.

Note the spelling: albumen, albuminous, albuminiferous, asbestos, gramme, radical, appara' tus (sing. and plu.) Fāte, fāt, fār, mēte, mět, pīue, pǐn, marîne, nōte, nŏt, möve, tūbe, tŭb, rūle, m \bar{y} , $\bar{y}=\bar{1}$.

PART II.—GLOSSARY.

ABBREVIATIONS.—AmOH for ammonia; BaCl₂ for barium chloride; CO₂ for carbonic acid: CnSO₄ for copper sulphate or blue-tone; FeSO₄ for copperas; HCl for muriatic or bydrochloric acid: HNO₃ for nitric acid: H₂O for water; H₂S for sulphuretted hydrogen; H₂SO₄ for sulphuric acid: KOH for caustic potash; NaCl for common salt: NaOH for caustic soda: Na₂CO₃ for carbonated soda: Na₂SO₄ Glauber salts; NH₄OH for ammonia: NH₄HS for ammonium sulphydrate; NO₂ for nitrogen oxide; SO₂ for sulphur dioxide.

Acid, Acetic.—CH_b.COOH, or C₂H₄O₂ (Germ.: Essigsāure; conc.: Eisessig; fr.: Acide acétique, vinaigre radical). Synonyms: Wood vinegar, wood acid, pyroligneous acid, tar acid, acetous acid, acidum aceticum. Apt impurities are sulphates, chlorides, lead, copper, zinc, iron, and lime. It ought to be colorless, and should evaporate entirely; the hydrometer is not a reliable mode of proof of strength. The determination of the value should be made by titrating with a standard alkali, using phenolphthalein as the indicator.

Acid. Carbolic.—C_tH_tOH, or C_tH_tO (Germ.: Carbolsäure, Phenylsäure; fr.: Acide carbolique). Synonyms: Phenol, phenic acid, phenyl alcohol; is noted for its antiseptic and disinfecting properties.

It has an extremely limited use in the dye-house, where it is sometimes employed to prevent decomposition. It is one of the products of coal-tar, and a dangerous poison.

Acid, Citric.— $C_bH_4(OH)(CO_2H)_3$, or $C_bH_5O_7$ (Germ.: Citronensāure; fr.: Acide citrique). Apt impurities are tartaric acid and sulphuric acid. Citric acid should evaporate clean.

Acid, Lactic.—C.H.(OH)CO.OH, or C.H.O. (Germ.: Milch-säure; fr.: Acide lactique). Apt impurities are chlorides, nitrates, sulphates, acetic acid, butyric acid. It is well to make an ash determination. Test should also be made for glycerine and glucose.

Acid, Muriatic.—HCl (Germ.: Salzsäure, Chlorwasserstoffsäure; fr.: Acide muriatique, chlorhydrique, hydrochlorique, esprit de sel marin, acide marin). Synonyms: Hydrochloric acid, spirits of salt, marine acid. Apt impurities are sulphuric acid, sulphurous acid (test by adding zinc and noting escape of sulphuretted hydrogen), free chlorine, ferric chloride (test with ammonia sulphocyanate), and arsenic. Common salt is sometimes added to raise the specific gravity of a weak acid. It is very readily detected by putting a little of the suspected acid in a saucer, and evaporating it. Sulphurous acid may be detected by a few particles of tin: if sulphurous acid were present, the offensive odor of sulphuretted hydrogen would be perceptible. Dilute is 1: 3.

Acid, Nitric.—HNO₃ or NO₂.OH. (Germ.: Salpetersäure; fr.: Acide nitrique, esprit de nitre). Synonyms: Aqua fortis, strong water, double aqua fortis. The coloration of ordinary nitric acid is due to the oxides of nitrogen. Apt impurities: Look for H₂SO₄ and HCl. The test for nitric acid is: Pour some H₂SO₄, conc. and e. p., upon the suspected substance in a dry test-tube; see that the mixture is cool, and then gradually add a concentrated solution of ferrous sulphate, so that it lies upon the sulphuric acid but does not mix up with it; a ring forming at the point of contact between the two layers is violet-red or brown, according to the amount of nitrate originally present. The reaction is:

10 FeSO₄ + 3H₂SO₄ + 2HNO₃ = 3Fe₂(SO₄)
$$_3$$
 + 4FeSO₄,
N₂O₂ + 4H₂O.

To detect free IINO₃ in the presence of a nitrate, evaporate the solution in a water bath with white wool, which will be turned to yellow. Dilute is 1: 2.

Acid, Nitrous.—HNO₂ or NO.OH. (Germ.: Salpetrige Säure; fr.: Acide nitreux, azoteux). To test for: Make a saturated solution of sulphanilic acid, and a saturated solution of naphtylamine hydrochloride. Mix the two solutions and add to the suspected liquid, when a brown color will show the presence of NO₂.

Griess's reagent for nitrous acid is a 0.5 per cent. solution of m-phenylenediamine with a small amount of sulphuric acid. If the solution was not colorless, it should be decolored with animal charcoal. The test yields a yellowish-brown color if traces of nitrous acid were present.

A solution of sulphanilic acid and naphtylamine sulphate is colored red by traces of nitrous acid. The solution to be tested is acidified with sulphuric acid, sulphanilic-acid solution added, and, after a few minutes, a small quantity of solution of naphtylamine sulphate decolorized by animal charcoal.

Acid, Oxalic.— $\begin{cases} CO.OH \end{cases}$ or $C_2O_2(OH)_2$, or $C_2H_2O_4$ (Germ.:

Oxalsäure; fr.: Acide oxalique). It can be obtained by the action of nitric acid upon sugar and starch, and is prepared on a large scale by treating sawdust or spent dye-woods with alkalies. Oxalic acid forms colorless transparent prismatic crystals, which have the specific gravity 1.64, are inodorous, intensely and unpleasantly sour, and do not grow moist on exposure. If they become damp, some nitric or sulphuric acid used in the preparation has not been thoroughly removed. It is soluble in its own weight of boiling water, but requires eight times its weight of water at 60° F. To detect sulphuric acid, dissolve in pure water, and add first pure hydrochloric acid, and then chloride of barium. If the oxalic acid is pure, the liquid will remain clear, but if there be an impurity of sulphuric acid, a white turbidity will appear. organic impurities, heat a portion with concentrated sulphuric acid. If any such matter be present, a small quantity of extract of indigo, boiled along with the solution of the sample, will have its color destroyed. If iron, lime, potash, or soda be present, a portion of the acid, heated to redness, will leave a fixed residue.

Acid, Pyroligneous.—(Germ.: Holzessig, Holzsüure; fr.: Acide ligneux, lignique, vinaigre de bois.) Symonyms: Black liquor, wood vinegar—is the crude acetic acid. Acetate of lime, when distilled with phosphoric acid, gives of the acetic acid in shape to be directly titrated with phenol-phthalein; the distillation had best be repeated three or four times.

Acid, Sulphuric.—H₂SO₄ or SO₂OH. (Germ.: Schwefel-OH.

säure; fr.: Acide sulfurique, sulfacide, huile de vitriol). Synonyms: Oil of vitriol, oil of sulphur. Manufacturers have adopted as "oil of vitriol" an acid which holds 93.5 per cent. H₂SO₄. Normal is 49 gr. per liter Its test is BaCl₂—heavy white precipitate. Impurities are: Nitrogen oxides, tested for by diluting somewhat, adding diphenylamine, which develops a blue; or nitrogen peroxide, by adding the acid, very much diluted, to a solution of potassic iodide and starch paste;

or cone. solutions of copperas and sulphuric acid. For calcium, iror, and arsenic, evaporate to dryness first; then, for iron, dissolve in nitric acid and add yellow prussiate of potash. For arsenic, use Marsh's test, or Reinsch's, which latter should be used in presence of SO₂ or NO₂. Examine for lead, which falls as a milky precipitate (PbSO₄) when the acid is diluted. Dilute sulphuric is 1: 5.

Acid, Sulphurous.—SO₂ 11 or H₂SO₃ (Germ.: Schweflige

Säure; fr.: Acide sulfurenx). A product of the combustion of sulphur. Cold water dissolves over 30 times its volume of sulphurous oxide; the solution contains hydrogen sulphite, or sulphurous acid, and may be kept unchanged so long as air is excluded. Excess of oxygen alters sulphurous into sulphuric acid.

Acid, Tannic.—(Germ.: Gerbsäure; fr.: Acide tannique.) The only reliable method of estimating tannic acid goes under the name of hide-powder method, and success depends upon the quality of the hide-powder.

Acid, Tartaric.— CH (OH). COOH: or C₄H₆O₆. (Germ.:

Weinsäure, Weinsteinsäure; fr.: Acide tartrique, tartarique). Synonym: Di-hydroxy-succinic acid. Impurities are lead, copper, iron, lime, sulphates, and bisulphate of potash. Good tartaric acid should evaporate white. The aqueous solution should not effervesce with nitric acid, as that would show carbonates. Test the nitric acid solution with baric chloride for sulphates, with silver nitrate for chlorides, and with ammonia and ammonium oxalate for lime.

Albumen.—(Germ.: Eiwciss, Eiwcissstoff; fr.: Albumine.) Precipitated by alcohol, corrosive sublimate (bichloride of mercury), solutions of lead, silver, and copper, acetic and muriatic acids, creosote, etc., but not by tannin, which latter will precipitate gelatine; thus discriminate between them. Solutions keep better when sodic arseniate is added. Turpentine oil is said to whiten dark-colored grades. Albumen is soluble in cold, but insoluble in hot water; at about 140° F. it coagulates. It takes the whites of ten dozen eggs to make a pound of albumen.

Blood Albumen has much the same properties as egg-albumen, but, unlike egg-albumen, is not coagulated by shaking

with ether. It occurs in blood-serum. Care must be taken in its manufacture, otherwise the red corpuscles will break and spoil the color of the product.

Alcohol.—(Germ.: Alkohol, if rectified, Weingeist; fr.: Alcool.) Ethyl-alcohol (C₂H₅.OH or C₂H₆O) is the ordinary alcohol of commerce. Synonyms: Spirit of wine, aqua vitæ, grain alcohol. Methyl-alcohol (CH₂.OH or CH₄O), or wood-alcohol, wood-naphtha, is a lower homologue. Apt impurities: Sugar, extractive matter, turpentine, coal-naphtha, pyroligneous and fusel oil. To detect the sugar, extractive matter, or turpentine, the best method is to distil a portion and take the specific gravity again. Fusel oil can be detected by its characteristic odor on rubbing some of the alcohol in the palms of the hands. Methyl-alcohol should not cloud on mixing with water, must remain colorless on addition of sodium hydrate, and when sulphuric acid is added it must turn, at most, only a weak yellow. Acetone should be looked for.

Alcohol, Aldehydes in.—(Germ.: Aldehyd; fr.: Aldehyde.) Guyon's reagent for: Dissolve 1 gr. fuchsine in 1 liter water, and add a mixture of 20 cc. sodium-bisulphite solution of 30 Bé. and 10 cc. concentrated hydrochloric acid. One cc. of this reagent added to 2 cc. of the solution to be tested will show the presence of aldehyde by the formation of an intensely purple-red reaction. This reagent is also known as "Schiff's reagent."

Alcohol, Acetone in.—Gunning's test for: Add to the solution to be tested tincture of iodine and ammonia; iodoform and a black precipitate of nitrogen iodide are formed, the latter gradually disappearing and the yellow color of the iodoform persisting. Alcohol by this treatment produces no iodoform.

Alkali. See Soda Ash.

Alkanet.—(Germ.: Alkannawurzel, Ochsenzungenwurzel; fr.: Oreanette.) The root of Anchusa tinctoria. Used for coloring ointments, cheeses, and pomades.

Alkermes. See Kermes.

Alligin.-Different species of plant glue.

Alum.—(Germ.: Alaun; fr.: Alun.) The difference between the different kinds of alums found in the trade will be best shown by giving a tabular view of three of the more ordinary alums found in the trade:

Common or	AMMONIA-	SULPHATE
Potasu-Alum.	ALUM.	OF ALUM.
$A1_2(SO_4)_3$	A12(SO4)3	
$+ K_2 SO_4$	+(NH ₄) ₂ SO ₄	
$+ 24 H_4 O.$	+ 24H ₂ O.	A12(SO4)3.
Alumina 10,8	11.4	15,5
Potassium 10.0	-	
Amonium	3.8	
Sulphuric Acid 33.8	35.3	36.0
Water 45.5	50.0	48.6

Al₂(SO₄)₃ + 18H₂O is "cake alum," "concentrated alum," or "patent alum": obtained from cryolite and bauxite. Iron is the most harmful impurity. A mixture of the ferrocyanide and ferricyanide of potassium may be added to the solution of a portion of the alum. An immediate blue precipitate shows the presence of iron. On longer standing, as, e.g., for an hour, this mixture may produce a blue precipitate even in pure alum. Tincture of galls, or a solution of tannin in any form, will produce a black color in the solution of alum, if iron be present. Alum gives acid reaction with litmus and phenol-phthalein, but not with methyl orange, unless some free acid is present. Alums of potash and ammonia are used interchangeably in industries; the soda-alum is extremely soluble in water, and for this reason it is about impossible to separate it from ferrous sulphate by crystallization.

Alum, Calcined.—Although good modern alum cannot be benefited by calcining, an inferior or an impure alum would be improved. The heat would have a tendency to render the iron in an impure alum insoluble in water by expelling a portion of the acid with which it was combined; if the alum was also of a very acid nature, some of the excess of acid would also be removed and its quality improved.

Amidon Grillé.-Synonym for brown dextrines.

Ammonia.—NH₄.OH, or AmOH, or NH₃.H₂O. (Germ.: Ammoniak; fr.: Alcali volatil, ammoniaque). Synonyms: Hartshorn, volatile alkali. Apt impurities are chlorides, carbonates, sulphides, iron or lead compounds, lime and tarry substances. Rub a little of the ammonia on the palm of the hand; if there were empyreumatic oil in the sample, the odor will betray it. The test can also be made by adding slowly a large excess of concentrated sulphuric acid, which chars the organic matter and gives a black color to the liquid. Ammonia should vola-

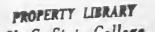
tilize completely, and not evolve a disagreeable odor on the addition of acid. The usual test for ammonia is the Nessler test. The reagent is made by dissolving 2 gr. potassic iodide in 5 cc. of water, warming and adding iodide of mercury as much as can be dissolved, cool and add 20 cc. H.O. let settle. and filter, and to 20 cc. of the liquid add 30 cc. of concentrated caustic soda c. p. A trace of ammonia produces with this reagent a vellowish-brown precipitate.

Ammonium Sulphide.-(NH4)2S or Am2S. (Germ.: Schwefelammonium, Ammoniumsulfid; fr.: Sulfure d'ammoniaque). Is prepared by saturating three parts of ammonia with H.S. diluting this solution with two parts of AmOH, when a sulphide is obtained which contains a little free ammonia.

Aniline Dyes .- (Germ .: Anilinfarben; fr.: Teintures d'aniline.) To test for dextrine, Glauber's salts, sugar, starch, sulphate of magnesia, and zinc oxid, dissolve in alcohol or ether, and observe amount insoluble-dextrine, starch, Glauber's salts. Dissolve in water and evaporate to dryness, when the erystals of Glauber's salts will form in large size. Dissolve in water and allow to settle, and after two or three days see how much deposition there has been. Steam the dve and observe if it gets sticky, indicating starch. Place in the sun or on the stove for a while; this will remove water of crystallization from Glauber's salts, leaving the crystals white, when they can be readily seen.

Mixtures of aniline dyes can be best shown by dusting a very little on to filter-paper which has been moistened. On account of the different rates of solubility, the components of the mixture will separate; a similar experiment can be made by dusting a little of the aniline dye on to a beaker nearly full of water. As the particles fall down through the water, they will separate and can be easily recognized.

Aniline Oil (Germ .: Anilinol; fr.: Huile d'aniline) should dissolve completely in hydrochloric acid or sulphuric acid, and the boiling point should not be changed by treatment with caustic soda. To determine the proportion of aniline and toluidine in an oil boiling between 185° and 205° F., take a known weight and treat it with one-half per cent. oxalic acid dissolved in water. Boil, and when dissolved cool down to 80° F., stirring continually. The greater part of the oxalate of toluidine precipitates: decant rapidly, throw the precip-



itate into a cheese-cloth, and squeeze out all excess. Then decompose this by ammonia containing enough alcohol to maintain in limpidity, heat, and when the solution cools the toluidine will crystallize out.

Annotto.—(Germ.: Orlean; fr.: Rocou, roucou, anotto.) Synonyms: Anotta, annatto, anoto, arnotto. Used in silk dyeing, and very rarely cotton, as it is a loose color. It is a pulp or paste, like putty, sometimes a solid cake, red inside, but brown on the exterior. Comes wrapped in large leaves. From seeds of bixa orellana, a South American shrub. In practice it is mixed with an alkali, since it is not soluble in water. Apt adulterants are brick-dust, red ochre, and iron oxide. The coloring principle is bixina, a red; another coloring matter is orelline, which is yellow.

Antichlor.—An agent for removing last traces of chlorine from goods. Bisulphite of soda, hyposulphite of soda, sulphurous acid, and ammonia have been used.

Antimonium.—A new antimony mordant, to replace tartar emetic.

Apparatine.—One of the numberless preparations of starch made by the treatment of flour and starch with eaustic or carbonated alkalis.

Aqua Fortis. See Nitric Acid.

Aqua Regia.—(Germ.: Königswasser; fr.: Acide azotique, can forte.) Three parts muriatic acid, and one part nitric acid. Synonyms: Nitromuriatic acid, azotic acid, nitrohydrochloric acid.

Arabil.-Different species of plant glue.

Archil.—(Germ.: Orseille, Färberflechte; fr.: Orseille.) Synonym: Orchil. This is a coloring principle prepared from a kind of moss, Roccella tinctoria, variolaria orcina, lichen corallenus, growing on rocks and stones, commonly called a lichen. The lichen has no color of itself, but by fermentation and treatment with lime and ammoniacal fluids, as urine, the coloring matter is developed. Archil has a peculiar characteristic smell, and was at one time used in silk and woolen dyeing to obtain different colors. Cudbear and litmus are very similar to archil. It is to-day almost entirely substituted by the aniline dyes.

$$\mathbf{Argol.-} \left\{ \begin{matrix} \mathrm{CH} \ (\mathrm{OH}). \ \mathrm{COOK} \\ \mid \\ \mathrm{CH} \ (\mathrm{OH}). \ \mathrm{COOH} \end{matrix} \right\} \quad \text{or} \ C_4H_4O_6KH \ (\mathrm{Germ.:} \quad \mathit{Roher} \right.$$

Weinstein; fr.: Tartre brut). When purified it is called "Cream of Tartar," or simply "Tartar" (Germ.: Weinsteinrahm, Cremor Tartari; fr.: crême de tartre). In its crude state it is either red or white argol, according as it has been deposited during the fermentation of red or white wine. Grapejuice, allowed to stand, deposits a crystalline layer upon the sides of the barrels. This is potassium bitartrate, and in commerce it is known under the name of argol. It is properly classified as one of the assistants in dyeing. The crystals, when shaken together, ring like fragments of earthenware. As it is relatively high-priced, many substitutes have been suggested, which generally consist of mixtures containing oxalic acid, bisulphate of potash, alum, common salt, etc. "Super-argol" is said to be a mixture of white argol and sulphuric acid. Apt adulterations of argol are organic matter, sand, carbonates, sulphates, chlorides, and lime. The latter from "plastered" wine. Look for starch also.

The best quality of argols is taken from the sides of barrels, and not from the bottom.

The sodium-potassium salt $(C_4H_4O_6KNa + 4H_2O)$ is known as Rochelle or Seignette Salt (Germ.: Seignette Salz; fr.: Sel de seignette), and used in medicine.

Arsenates or Arseniates.—(Germ.: Arsensaure Salze; fr.: Arseniates.) Arsenate of Potash.—K₂HAsO₄ + 12H₂O. (Germ.: Arsensaures Natrium; fr.: Arseniate de potasse). Formerly used as a resist in combination with pipe-clay.

Arsenate of Soda.—Na₂HAsO₄ + 12H₂O (Germ.: Arsensaures Natrium; fr.: Arseniate de soude). This product is largely used in calico-printing as a dung-substitute in clearing the cloth after mordanting, to remove the excess of unfixed mordant.

Arsenic Tests.—Some simple tests are: A gas-jet is turned down to quite a pin-point, until the flame is wholly blue; then the suspected sample is brought into contact with the outer edge of the gas flame, and a bluish-white coloration, due to arsenic, will be seen in the flame (test No. 1). The fumes that are given off will be found to have a strong garlic-like odor, due to the vapor of arsenic acid (test No. 2).

Another test for arsenic is Gutzeit's test: The substance to

be tested, together with c. p. zinc and c. p. dilute sulphuric acid, is placed in a test-tube, the mouth of which is closed with a piece of filter-paper moistened with a few drops of a silver-nitrate solution. The arseniuretted hydrogen formed if arsenic is present produces a yellow spot on the paper, which is blackened upon moistening with water.

The usual method of testing for arsenic is Marsh's test, which is made by developing arseniuretted hydrogen; this is burned and a piece of cold porcelain is inserted in the flame,

when a metallic mirror deposits on the plate.

Antidotes in Cases of Arsenic Poisoning.—Emetics, such as warm mustard-water (one teaspoonful of mustard in a glass of water), or a quantity of soap-suds. Also milk, the whites of eggs, or freshly precipitated ferric hydrate ($\mathrm{Fe_2H_6O_6}$).

Bark (See Quercitron Bark).—The term bark is generally used to designate quercitron bark. The barks from other trees have been used as dyes, but the word "bark" is almost exclusively limited to quercitron. Alder bark, oak bark, pomegranate bark, pine bark, and willow bark are all historical

dyes, and cut no figure in practical applications.

Barwood.—A red dyewood from Africa, closely resembling sandal-wood. Its coloring-matter is not easily extracted by water, for boiling only dissolves a small quantity of it, and this precipitates in great part as the water cools; there is, therefore, no barwood liquor or extract. To dye with it, the rasped or ground wood has to be used just as madder is used in madder-dyeing. The goods take the color from the water as fast as it takes it from the wood; the coloring-matter is gradually transferred until the desired shade is obtained or the wood spent. It produces a good imitation of the Bandana Red, but it requires a very skillful dyer to get a good result. It was used for the production of "mock turkey red."

Benzene.—C₆H₆. Synonyms: Benzol, coal-tar naphtha. It is better to use the name benzol, as that distinguishes it clearly from benzine, a product from the petroleum industry.

Berries.—(Germ.: Gelbbeeren, Avignonkörner; fr.: Graines d'Arignon.) Synonyms: Persian berries, French berries. There are about seven or eight different qualities, all derived from the same kind of shrub, Rhamus infectoria, in France called the dyer's buckthorn; it grows in France, the island of Candia,

Wallachia, and Asia Minor. One variety grows in Persia, whence its name.

Bicextine.—A sizing material made by malt process to a dextrin.

Bichrome or Bichromate of Potash or Soda.—Synonym: Chrome. Rarely sophisticated.

Bichrome of Potash.—K₂Cr₂O₇ (Germ.: Kaliumbichromat; fr.: Bichromate de potasse).

Bichrome of Soda.—Na₂Cr₂O₇ (Germ.: Natriumbichromat; fr.: Bichromate de soude).

Bichrome, used for mordanting, should be 3 per cent., calculating on the weight of the goods, and tartar, 2 per cent. Sulphate of copper, 2 to 3 per cent., is advantageous. Apt impurities: Sulphates.

Blanc Fixe.—Trade name for barium sulphate. Used in sizings for weighting and filling the goods.

Bleaching Powder.—Synonym: Chlorinated lime. U. S. P. Method for Estimating Chlorinated Lime.—If 0.354 gramme of chlorinated lime be thoroughly triturated with 50 cc. of water and carefully transferred, together with the washings, into a flask, and then 0.8 gr. of iodide of potassium and 5 cc. of diluted hydrochloric acid added, the reddish-brown liquid, mixed toward the end of the titration with a few drops of starch, test solution, should require, for complete decoloration, not less than 35 cc. of decinormal sodium hyposulphite solution, and each cc. corresponds to 1 per cent. of available chlorine.

Blue Stone. See Copper Sulphate.

Borax, Borate of Soda.—Na₂B₄O₇ (Germ.: Tetra (saures) borsaures Natron, Tinkal; fr.: Borax, biborate de soude, soude boratée). A few years ago the main source was Tuscany, but now it is brought in large quantities from Nevada. It is little used in dyeing, but is used more or less in the woolen mills in connection with oil, in order to make emulsion for oiling the wool as it stands before the picker. "French borax powder" was a mixture of soda ash and lime.

Brazil Wood.—(Germ.: Fernambuk, Fernambukholz, Brasilienholz, Rothholz; fr.: Bois de Brésil, Bois de Fernambuk, brésillet rouge de Brésil.) Synonyms: Brazil, Pernambucowood, Red-dye-wood. This is a red wood, Casalpina crista, and, as its name indicates, comes from Brazil. It can be distinguished from barwood by the fact that it speedily gives a

bright-red color to water. Sandal-wood and barwood do not, under similar circumstances, color water. Apt adulterations are common salt and nitrate of soda. Liquor should be heated with some nitric acid until the color is destroyed, then the residual liquor should be mixed with distilled water and a solution of silver nitrate should be added. It is from the same kind of tree and nearly identical with peachwood, Limawood, and sapan-wood. The richest variety is from Pernambuco, and is sometimes called Pernambuco-wood. Owing to the fugitive character of the color, it is being less and less used.

Brillantkleister .- Different species of plant glue.

Camwood.—(Germ.: Cambalholz, Angolaholz, afrikanisches Rothholz; fr.: Bois de chum, bois d'Angolc.) Synonyms: Kambewood, Angola-wood. This is the wood of Baphia nitida, obtained from the west coast of Africa, and has about the same properties as Brazil-wood. Like other dye-woods, it is steadily decreasing in application.

Canaigre.—A tuberous root (Rumex hymcnoscepalus) being increasingly cultivated in the Southwestern parts of the United States and in Mexico, and used for the tannin which it contains.

Caseine.—(Germ.: Casein, Käsestoff; fr.: Caseine, caseum.) Curd milk, sold as a powder under the name of lactarine. It is prepared by precipitating skimmed milk with dilute acids. It is insoluble in pure water, and must be dissolved by solutions of ammonia and borax, the latter being preferred on account of its antiseptic properties. It is used in calicoprinting as a substitute for albumen. It is not coagulated by steaming, and can only serve for colors requiring a moderate degree of fastness. It is distinguished from albumen by the circumstance that it does not become insoluble when heated in diluted solutions.

Catechu.—(Germ.: Catechu, japanische Erde; fr.: Cachou. terre du Japon.) Synonyms: Cutch, Japan earth, terra Japonica, kut, cashew. Bombay catechu is prepared from the wood and fruit of the catechu palm, Arcca catechu. Bengal catechu is made from the twigs and unripe pods of Mimosa catechu. Gambier catechu, also called Cubical and Yellow Catechu, is obtained from the leaves of the shrub Uncaria gambier, and occurs in commerce in the form of small cubes.

Good catechu should not contain more than 5 per cent. ash, or more than 12 per cent. of matter insoluble in boiling alcohol.

Its texture is resinous, and, if good, sufficiently brittle to break under the hammer. The color, however dark outwardly, should be a brownish cream color within. If it is deep-brown throughout, and soft and pitchy in consistence, so as to cling to a knife, it has become impaired in quality either by long keeping or by exposure to moisture. It should contain about half its weight of tannin, varying in this point from 37 to 56 per cent.

Catechu is entirely soluble in hot water, if genuine. The brown solution, when cold, lets fall a sediment, containing the bulk of the catechuic acid.

Catechu is frequently adulterated with clay, sand, starch, ochre. etc. The detection of these impurities is very easy. The sample is boiled in alcohol and the decoction strained, when all such admixtures remain and may be dried and weighed. Blood or sugar may be detected by burning the admixture, when the characteristic odor betrays them.

Chemic.—Synonyms: Bleaching powder (which see), chloride of lime, and bleach. Bleaching powder is now made in this country with as high a per cent. of chlorine strength as the best grades of imported chemic.

Chinese Green (Lo-kao, Vert-Venus).—Prepared from the buckthorn, superseded by the aniline greens,

Clay.—(Germ.: Thon, Lehm; fr.: Argile, glaise.) Synonyms: Terra alba, kaolin, porcelain clay, China-clay, pipeclay, Devonshire-clay, Cornish-clay, Argil. Sometimes used by the printers as resists, and they are abused for adulterating lake-colors and other pigments. China-clay is an hydrated silicate of alumina, and is obtained from orthoclase or felspar. It differs from porcelain in not possessing such strong affinity for water, which difference may be readily shown by testing with the tongue. The porcelain-clay absorbs water from the tongue and adheres to it, while China-clay will not. The tests are as follows:

1st. It must not grit between the teeth.

2d. It must have an unctuous feel, and should be compared with a standard by putting on a glass plate, adding a little water and rubbing each lot, the new and the old.

3d. Should not effervesce when muriatic acid is added.

4th. Color should be compared to standard.

5th. Avoid iron, which would alter the color. When boiling, add soap with tallow in order to check spattering.

Chlorophyll.—(Germ.: Chlorophyll, Blattgrün; fr.: Chlorophylle, vert des feuilles.) Synonym: Leaf-green. The green coloring-matter of the leaves. Insoluble in water. In the attempt made to use it as a dye, grass has been first boiled out in water, and the color extracted from the residue by a very weak lye of carbonate of soda, from which the chlorophyll is thrown down as a paste by the cautious addition of an acid. It has been experimentally used in dyeing and printing, but not with satisfactory results, as it is dull, fugitive, and very low in tinctorial power, and consequently expensive. Xanthophyll and Erythrophyll are yellow and red colors found in decaying leaves. They are of no practical importance.

Cochineal.—(Germ.: Cochenille, Johannisblut; see Kermes; fr.: Cochenille.) An insect, coccus polonicus, feeding on a species of cactus in Central America, cactus opuntia. The usual adulteration consists in adding some powdery matter to the cochineal, to increase its weight. French chalk, white lead, and ground tale are said to be chiefly used. These adulterations are easily detected upon analysis, for by calcining the mixture at a red heat, the true matter of the cochineal is burned away, leaving the added mineral adulteration, the quantity and nature of which can be then ascertained. The pure coloring-matter of cochineal has received the name of carmine.

Copperas.—FeSO₄ + 7H₂O (Germ.: Grüner Vitriol, Eisenvitriol; fr.: Vitriol vert, couperose verte). Synonym: Green vitriol. A name given to the protosulphate of iron, from the mistaken notion that it contains copper. If of a dull, whitish-green color—as it is technically called, milky—the presence of alumina is to be suspected. Add pure caustic soda—that prepared from metallic sodium to be preferred—in large excess; boil in a clean iron vessel and filter. Add to the clear filtrate a solution of pure sal ammoniac. If, on standing, a white precipitate appears, alumina was present in the copperas.

Lime is sometimes dusted over copperas to give that brown spotty appearance which some consumers prefer.

The direct uses of copperas in dyeing have very much diminished. For dyeing blacks upon wool in conjunction with

logwood it has been, to a very great extent, superseded by chrome. For blacks upon cotton, as also for saddening drabs, clarets, etc., the nitrate of iron is generally preferred.

Copper Sub-Acetate.— $Cu(C_2H_2O_2)_2 + CuH_2O_2$ (basic copper acetate) (Germ.: *Grünspan*; fr.: *Verdet*, *vert-de-gris*). Synonyms: Verdigris and verditer. Apt impurities are chalk and sulphate of copper. Very poisonous. Antidote: Whites of eggs. milk, magnesia.

Copper Sulphate.—CuSO₄ + 5H₂O (Germ.: Kupfervitriol, blauer Gallitzenstein; fr.: Vitriol de cuivre, vitriol bleu, vitriol de Chypre). Synonyms: Bluestone, Cyprus, Roman vitriol, blue vitriol, vitriol of copper, copper-vitriol. If it holds FeSO₄ it is called Admont, mixed, Bayreuth (or Salzburg) vitriol. Admont vitriol: 5 parts FeSO₄ and 1 part CuSO₄. Double Admont: 4: 1; Bayreuth, 7: 1; Salzburger, 17: 5% (also called Double Eagle vitriol).

Look for iron by boiling with a few drops HNO₁, and then adding AmOH to dissolve the precipitated oxide of copper, which leaves then the iron behind as insoluble oxide. Look for zinc by first removing the iron by AmOH, and add (NH₄)HS.

Cudbear.—(Germ.: Persio; fr.: Orseille de terre épurée, orseille violette.) A modified extract of colorable lichens (Lecanora tartarea, Lychen omphalodes, calcareus, saratilis), similar in its behavior to archil. Its peculiar odor is easily remembered. Its use is confined to a few cases of silk dyeing. It is sometimes purposely contaminated with mineral matter, such as salt, carbonate of lime, etc. These frauds may be detected by burning a weighed quantity of the sample to ashes.

Cutch. See Catechu.—The test to distinguish between cutch and gambine is to dissolve in potash, and extract with benzine, when there will be a green fluorescence if gambine is present.

Delaine.—(Germ.: Wollmousseline; fr.: Muslin de laine, mousseline de laine.) Although these words in French distinctly indicate a fabric made of wool, they are employed in English to indicate a material of which the weft only is wool, the warp being cotton. In French this fabric is called chaine cotton—that is, "cotton warp"; and sometimes mi-laine—that is, "half wool."

Dextrine.—C_cH_nO_c (Germ.: Dextrin, geröstete Stärke, Stärke-gummi; fr.: Dextrine). Synonyms: British gum, artificial

gum, torrified starch, starch-gum, Leiogomme. Apt impurities: Unconverted starch (tested for with iodine), grape-sugar, acids. Adulterations may be sand, plaster-of-Paris, sulphate of barium, and tale. The dextrine is dissolved in water, evaporated to a syrup, and then precipitated by alcohol. The glucose can be estimated by Fehling's test, but must be boiled only for a short time. Comparative experiments showing thickening power should be made. A quick method for testing is to mix it with cold water in a long test-tube; in the course of a few hours all the insoluble or imperfectly calcined and raw particles fall to the bottom of the test-tube and from their quantity the quality of the sample may be judged. Test for grit: If the lumps yield to pressure the residue is most likely charred particles of farina. It is one of the characteristics in dextrine that, when hard, it becomes thin, and grit will sink to the bottom; other gums are apt to keep the grit in suspension.

Brown Dextrines.—Synonyms: Amidon grillé, gebrannte Stärke, Gomein, Gommeline, Lefèbregummi, Leiocomme, Leiogomme.

Diastase.—(Germ.: *Diastase*; fr.: *Diastase*). A peculiar substance contained in malted grain, and is used for removing starch sizing out of cloth, where a soft finish is demanded. One part of this active ferment is sufficient to change 2,000 parts of starch into dextrine and sugar.

Divi-Divi.—(Germ.: Dividiri; fr.: Diridivi.) Synonym: Libi-divi. This is an astringent substance from pods, Poinciana coriaria, possessing some of the properties of sumac; attempts were made to extend its use in dyeing, but not with much success; it is used by tanners, but scarcely, if at all, by the dyers or printers.

Egalisol.—This is borosulphate of soda. Three per cent. of egalisol is used with 4 per cent. of bichromate to even up the color.

Emeraldine, made from aniline and potassium chlorate.

Epsom Salts.—MgSO₄ + 7H₂O (Germ.: Bittersalz, Englisches Salz, Schwefelsaure Magnesia; fr.: Sel d'Epsom, epsomite, mugnésie sulfatée, sel amer). Synonym: Bitter salt. Used in certain finishes.

Erythrophyll. See Chlorophyll.

Farina.-(Germ.: Kartoffelstürke; fr.: Fécule de pommes

de terre.) Synonym: Potato-flour. This term is usually applied in trade to the starch obtained from potatoes, often called potato-starch.

Fehling's Solution.—This consists of two solutions, namely (a) an aqueous solution of copper vitriol, which contains in one liter 34.639 gr. crystallized copper sulphate, and (b) a solution which is prepared by dissolving in a liter flask 173 gr. sodium potassium tartrate, adding 572 gr. caustic soda of 1.12 sp. gr. (containing 60 gr. sodium hydroxide), and filling up to the mark.

Flavine.—The yellow or orange color obtained from quercitron bark. Apt adulteration is common salt.

Flour.—(Germ.: Mehl; fr.: Farina.) Chemically, flour may be considered a mixture of starch and a peculiar substance called gluten. It is the amount of starch that renders it more or less valuable. The gluten has no sensible thickening power of itself. The best and only test of any practical value is to make a trial of it in thickening, observing how much it takes to give a good consistent paste.

The principal adulterations are potato-starch and beanflour, Indian-corn flour, rye- and rice-flour. Sometimes alum, chalk, bone-dust, and plaster. Moisture should be determined by drying in a water-bath until constant weight is obtained.

To detect potato-flour use a microscope. The particles of potato-starch are of various sizes; and, moreover, upon placing a little of the suspected flour before the microscope and moistening it with a solution of potash, the particles of wheatflour undergo no change, while the granules of potato-starch spread into thin transparent plates. Test for alum with logwood, and for mineral matter by shaking with chloroform, and allowing to settle: the flour will float.

Flour Testing.—Soak 5 gr. of flour in 100 cc. cold H₂O, for a couple of hours with frequent stirring; toward night put into a filter; fill up the funnel with H₂O, and let it stand through the night. Residue holds cellulose, starch, gluten, and fat; the filtrate-holds albumen, gum, and sugar. Boil the residue in 500 cc. H₂O containing 5 cc. H₂SO₆, conc. until the iodine test is not given; then neutralize with NaOH solid, and titrate against 10 cc. Fehling's solution in 40 cc. H₂O at 100° Centigrade, and report as starch per 5 gr. Boil the solution and divide into two parts. The first half is titrated against

2 cc. of Fehling's solution in 40 cc. $\rm H_2O$ at 100° Centigrade, and report as sugar per $2\frac{1}{2}$ gr. The second half is acidified with $\rm H_2SO_4$ dil., boiled until the iodine test is not given, neutralized with NaOH, and titrated against Fehling's solution without the addition of $\rm H_2O$. Excess is cane-sugar, which is to be reported as cane-sugar per $2\frac{1}{2}$ gr.

If flour be mixed and kneaded in a cloth with water, a whitish liquid will be strained off, which, being left to settle, will deposit starch. The substance retained by the cloth is gluten, which is white and insoluble. A poor flour is merely slimy.

Special names for flour paste boiled with alkalies: Poliokolle, Schlichtepräparat, Apparatin, Gallerte, Krystall-Appretur, Pflanzenleim, Alligin.

Fuller's Earth.—(Germ.: Walkerde; fr.: Terre à foulon, glaise à dégraisser, argile smectique.) A term originally applied to a lime or argillaceous earth occurring in association with chalk and oölite.

Fustet.—(Germ.: Fisetholz, Visetholz, ungarisches Gelbholz; fr.: Fustet.) Synonyms: Young fustic, Zante-fustic. Wood of a European tree, Rhus cotinus, obtained its prefix "young" on account of the smallness of its branches compared with that of the yellow wood, which was distinguished as "old" fustic. Its colors are more fugitive. Fustet is very little used in cotton-dyeing, not at all in calico-printing, but is used by some wool-dyers to give a more fiery tint to their scarlets.

Fustic.—(Germ.: Gelbholz; Bois jaune.) Synonyms: Cubawood, old fustic, yellow wood. The wood of a tree (Morus tinetoria) formerly known as "Dyer's Mulberry." The colors it gives are not very stable. It comes into sale in four states; namely, as chips, powder, aqueous extract, and as a paste or lake.

In the two former states it is generally laid up for several weeks before coming into use, being frequently turned over and sprinkled with water. This process softens the woody fibre, and enables the color to be more easily extracted. Sometimes, however, the water is present in such amount as to constitute an adulteration. Samples both of chipped and rasped fustic, carefully dried at 212° F., have been known to lose not less than 48 per cent.

Fustic liquor, or extract of fustic, is water saturated at

212° F. with the coloring principles of fustic, and boiled down to 5° or 10° Twaddle. Like the other extracts, this liquor is often "sprung." as it is technically called; that is, mixed with some substance that may increase its weight, and cause it to mark a higher degree on the hydrometer. Common salt is generally selected on account of its cheapness, ready solubility, and from the fact that it does not very seriously modify the appearance of the liquid. Look for quercitron, chestnut, curcuma, starch, dextrine, molasses, glycerine, zinc salts, anilines, and alizarines.

Scheme for Fustics.

Report No.

Date.

Laboratory No.

Their Mark or No.

Ash (mineral):

Water:

Specific Gravity:

Beaumé:

Twaddle:

Insolubles:

Alkali:

Alizarines:

Resins:

Glucose:

Gall Nuts.—(Germ.: Galläpfel, schwarze oder grüne; fr.: Noix de galle, galles noires-tertes.) Synonyms: Nutgalls, oak apples. This valuable dyeing material is an excrescence from a certain tree (Quercus infectoria): it is caused originally by the puncture of a little insect (Cynips quercus folii, Diplolepis galla tinctoria) on the leaves, or small branches of the tree, in order to deposit its egg in the cavity formed. The juices of the tree collect round the egg, and, hardening, form the gall nut. If the larvæ have eaten out, there is a loss of astringent principle, and the nuts are then called green galls.

Gambier resembles catechu or cutch. It is obtained by extracting the leaves of the *Uncaria gambier*, a shrub growing abundantly in the Malage and India Islands. It contains varying amounts of a tannin known as catechu-tannic acid or mimotannic, and a white crystalline body called catechine or catechine acid. Apt impurities are clay and occasionally starch. Excess of mineral is determined by igniting a weighed amount in a crucible. A good cutch should not leave more than 3 per cent. of an ash. The percentage of water varies

from 15 to 25. The estimation of the amount of catechu-tannic acid is said to be of no value.

Garanceux.—A preparation of madder. The distinction is that while garancine is prepared from fresh madder, garanceux is made from the spent or exhausted madder. It is weaker than garancine.

Garancine.—(Germ.: Garancin; fr.: Garancine, charbon sulfurique.) A formerly important red-dyeing preparation of madder, made by treating madder-root with sulphuric acid.

Gebrannte Stärke.-Synonym for Brown Dextrines.

Gelatine.—(Germ.: Gelatine; fr.: Gelatine.) Gelatine, when pure, is colorless, tasteless, inodorous, soluble in boiling water, but insoluble in cold, though it swells up and softens. It is insoluble in alcohol.

Glauber's Salts.—Na₂SO₄ (Germ.: Glaubersalz, Wundersalz, schwefelsaures Natron; fr.: Sel de Glauber, sel admirable, soude sulfatée). Synonyms: Sodium sulphate, wonderful salt, Sally Nixon, in corruption of the ancient name sal enixum. Sulphate of soda crystals (Na₂SO₄ + 10H₂O) should be neutral, or at most they should have a very feeble alkaline reaction. They should dissolve in water without leaving any residue, and, above all, they should be free from compounds of iron. Look for aluminum, magnesium, common salt, insolubles.

Glauber's salts is used almost universally in dyeing wool in acid bath to aid in making the dye go on even. The desiceated Glauber's salts is the more economical.

Gloriagummi.—Different species of plant glue.

Glycerine.—
$$C_3H_5O_3$$
 or C_2H_5 $\begin{cases} OH \\ OH \\ OH \end{cases}$ (Germ.: Glycerin, Ölsüss;

fr.: Glycérine). Synonym: Sweet principle of oils. It should evaporate without residue. Heating with sodium hydroxide should not color it; if it does, look for starch or sugar.

Gomein.—Synonym for Brown Dextrines.

Gommeline.-Synonym for Brown Dextrines.

Gum.—(Germ.: *Gummi*; fr.: *Gomme*.) Divided into gum proper and gum resin. The first being soluble in water, and the second substances as copal.

Gum-Arabic.—(Germ.: Arabisehes Gummi; fr.: Gomme arabique.) This is the gum of Aeaeia vera, nilatica, arabica. It can be taken as a type of gum proper, and next to it comes gum-senegal. It has a tendency to get sour, but this may be

partially corrected by the addition of soda. Next in order of value comes East India gum, sometimes called Turkey gum. Adulterations: Cheap and useless gums, also dextrine, starch. tragacanth, ceresin, and insolubles. A comparative test with an accepted standard in the matter of sp. gr. is of great value.

Gum-Senegal.—(Germ.: Senegalgummi; fr.: Gomme Sénégal.) This is the gum of Acacia Senegal, and differs from gumarabic chiefly by its darker color and larger amount of impurities, such as sand, earth, fragments of wood, etc.

Gum-Tragacanth (gum of Astralagus tragacantha).—This gum, when sold in lumps, is rarely adulterated, but in the state of powder it is often mixed with gum-Senegal. This may be detected by dissolving a portion of the suspected sample in hot water, and adding, under constant stirring, a few drops of the tincture of guaiacum. If gum-Arabic or gum-Senegal be present, a blue color will appear.

The subjoined table will be found useful in discriminating the various kinds of gums:

Gums.	Sulphate of Iron.	Tincture of Guiacum.	Subscetate of Lead.
Gum Arabic	Yellow Precipitate	Blue Color	White Curd
Senegal	Yellow Precipitate	Blue Color	White Curd
	Yellow Precipitate	Blue Color	Transparent Jelly
Tragacanth	Yellow Precipitate	No change	Transparent Jelly
Dextrine	No precipitate		

Gummi Germanicum.—Different species of plant glue. Gummi Saxonium.—Different species of plant glue.

Haï-Thao, or Gelose.—Algæ from Cochin China and the Mauritius.

Hemlock Bark.—(Germ.: Schierlingstanne; fr.: Sapin de Virginie.) The bark of the hemlock spruce (Pinus canadensis), a tree very plentiful in the northeastern States of the American Union.

It is, like sumac, an astringent, and has not been used for dyeing purposes with satisfactory results, as it gives a rusty or "foxy" surface reflection to the goods.

Hypernic.—A name given to Nicaragua-wood, and sometimes to any other red-wood, or red-wood extract of the same class. Hydrofugicolle.-Different species of plant glue.

Indigo.—(Germ.: Indigo; fr.: Indigo.) Obtained from the fermented and treated sap, essentially an East Indian product. The chief indigo yielding plants are various species of Indigofera—tinctoria, disperma, Agil, argentea, glauca, cœrulea, cineria, glabra, etc.

Artificial Indigo has now largely superseded the natural product; this is one of the triumphs of modern chemistry.

Indigo-Carmine or Indigo-Paste.—(Germ.: Indigocarmin, blauer Carmin, präcipitirter Indigo; fr.: Indigo soluble.) Synonyms: Blue carmine, soluble indigo, precipitated indigo, distilled blue, extract of indigo, chemic blue, Saxony blue.

The analysis should embrace the moisture, sp. gr., insolubles, dyeings on mordanted and unmordanted cotton, as well as gun-cotton, wool mordanted and unmordanted, and a dyeing as compared with the type. The reduction should be clear and even. Fehling's solution should not be precipitated. Test filtrate for starch, chlorides, sulphates, anilines. Examine the ash of filtrate. Look for dyewoods by oxalic acid and filter paper. Adulteration with coal-tar colors is proved by dyeing gun-cotton with the suspected sample; indigo alone will not dye the gun-cotton, but the basic coal-tar colors will. The odor should be noted; the per cent. of water should be found; boiling with sulphuric acid should not turn the paste black, indicative of organic matter; precipitated with salt, and filtered. should give a clear filtrate. This filtrate, when boiled, should give no odor of dextrine. The filtrate tested with iodine should not give a coloration (starch). Digested with sulphurie acid and then tested with Fehling's solution would show any sugar which had been inverted.

The laboratory method of reporting indigoes is as follows:

Report No.	their mark of No
Moisture	3.15
Ash	12.10
Specific Gravity	1.485
Indigotin and Indigo Red (1st deter-	
mination)	44.54
Indigotin and Indigo Red (2d deter-	
mination)	44.24
	$2)\overline{\underline{88.78}}$
Average of two determinations	44.39

Gross frauds, which are said to be sometimes attempted by the indigo makers, could be easily detected by chemical analysis, such as the admixture of ground slate, black sand, plumbago, lead-powder, starch, etc.

Indigo Sulphate, To make a Pure Extract of.—Dissolve crude sulphate of indigo in water, nearly boiling, and a quantity of old but clean white flannel or other woolen articles worked in it until saturated with color; then wash well in cold, and afterward in warm, water until the color begins to "bleed," that is, until the washing water begins to remove the blue and become tinged with it; they are theu treated with hot water containing a feeble proportion of carbonate of soda; this removes the blue color very rapidly from the woolen rags, leaving them of a dull brown color; the blue thus dissolved is considered as purified. A little acid added to the extracted blue enables it to dye up a good clear blue.

Irish Moss.—(Germ.: Irländisches (Perl) Moos, Carragaheen Moos; fr.: Chondre erispé; Chondrus (Fucus) erispus.) Synonyms: Pearl moss, Carrigeen moss, Caragheen moss. Contains a large proportion of a gelatinous matter, pectine, or vegetable jelly.

Iron Acetate.—Fe(C2H3O2)2 (Germ.: Essigsaures Eisenoxydul, Holzessigsaures Eisen, Essigsaure Eisenbrühe, Eisenbeize, schwurze Brühe, schwarze Küpe; fr.: Bouillon noir, tonne au noir, gris de fer, couleur de fer, acétate (de protoxyde) de fer). Synonyms: Ferrous acetate, black liquor, black mordant, pyrolignite of iron, printer's iron liquor, iron mordant. important mordant consists of protoxide of iron, combined with and kept in solution by crude wood-vinegar, or pyroligneous acid. It is prepared by two distinct methods: Either scrap-iron dissolved in the free acid-generally with the aid of heat-or a solution of copperas is mixed with a solution of some soluble acetate, such as acetate of lime, or lead; this is Chamoisbeize. Contrary to what might be expected, pure acid or pure soluble acetates do not produce as good a quality of Black Liquor as such containing a certain amount of tarry matter. Concerning the part played by this tar in dyeing, there is some diversity of opinion. It is found that the purified acid produces a less satisfactory liquor, a fact due, according to Moyret, to the presence of a small quantity of pyrocatechol in the crude acid, which forms a compound with the ferrosoferric oxide in the solution, and causes its intense color and keeping properties.

The liquor used by dyers is often concentrated, not by evaporation, but by the addition of copperas (ferrous sulphate), thus, the addition to 1 gallon of Black Liquor, sp. gr. 1.085, of ½ pound of copperas would raise its density to 1.111. Over 1 pound of copperas has been found in 1 gallon of iron liquor. Tannin also is sometimes added. Black Liquor is also prepared by the action of ferrous sulphate or acetate of lime; the liquor produced has an average density of 1.11, and always contains sulphate of lime.

Black Liquor is sold at different strengths, ranging from 10° to near 30° Twaddle. It has an astringent taste, is of an olive color, but in large quantities it appears black.

It serves for producing upon cotton a variety of shades according to its strength, and the coloring-matter with which it is used. It gives its oxide of iron to the fiber more readily than copperas.

The peracetate of iron-acetate, or pyrolignite of the peroxide, has been proposed as preferable to nitrate of iron, but has searcely received any other than mere experimental application.

To find if Black Liquor is genuine, add a little nitric acid, and apply heat to peroxidize the iron. Then add excess of ammonia, and filter off the precipitate. The clear liquid remaining is evaporated to dryness, and the residue heated strongly. Nothing should remain but a trace of earbon derived from tarry matter. If common salt had been added to raise the Twaddle, it will be found as a white mass.

Iron Nitrate.—Fe₂ (NO_3)₆ (Germ.: Salpetersaures Eisenoxyd; fr.: Rouille, pernitrate de fer). Synonyms: Iron nitrate " true," iron-mordant, rust-mordant. Made by dissolving iron in nitric acid; is made also by running concentrated nitric acid into a solution of copperas made by driving steam into the solid crystals of FeSO₄.

Iron for blue should be sharper than iron for other purposes; if too "dead"—that is, if the amount of iron be too great in proportion to the acid, a part of the Prussian-blue formed will be deposited at the bottom of the dye-pan, and that which is fixed upon the goods will be dull, loose, and

cloudy. The nitrate of iron must not be acid, otherwise the color is thinned and probably irregular.

Add to a small portion of the sample pure ammonia, till all the oxide of iron has been precipitated. This is then filtered off, and the clear liquor evaporated down to dryness and the residue heated to redness in a small porcelain capsule. If any fixed matter remains, the sample has been got up with nitrate of soda. Alumina is an objectionable impurity.

To ascertain whether any portion of the iron is in a state of protoxide, drop in cautiously a solution of carbonate of soda. If the sample be a pure peroxide, the precipitate will be of uniform pale yellow. If any protoxide be present, clouds of a greenish color appear in the liquid. Or a dilute solution of the red prussiate of potash may be added, which, if any protoxide exist in the sample, will give a blue precipitate.

Iron Protonitrate.—Fe(NO₃)₂ (Germ.: Salpetersaures Eisenoxydul; fr.: Protonitrate de fer). Made by combining 1,750 parts copperas with 2,080 parts nitrate of lead, both in solution; is also made by dissolving FeS in cold HNO₃ diluted.

Iron Salts.—A convenient test is a solution of tannin and a solution of logwood.

Iron, Sulphate of. See Copperas.

Japanleim. Different species of plant glue.

Johannisblut, Deutsche Cochenille (St. John's blood).—An old name for kermes, so named because it was collected with considerable religious ceremony on St. John's day. See Kermes.

Kermes.—(Germ.: Kermes; fr.: Kermès, Alkermes.) Synonym: St. John's blood, since it was collected from the oaktrees on St. John's Day with considerable religious ceremony. It is derived from the female insect Coccus Ilicis, which is found principally upon the Ilex oak (Quercus Ilex). At one time a great deal of kermes was sent to Venice to produce the scarlet to which city it owes its name.

Lac Dye.—(Germ.: Färbelack; fr.: Laque en teinture, laque en trochisques.) A color of the same class as cochineal. Consists of small parasitical insects—Coccus ficus (Laccw)—cemented together by a resinous matter exuding from the twigs of the trees which they inhabit. A good lac should be soft, so as to be broken by the fingers. The fracture should not have a shining resinous appearance, and should give out a

strong peculiar odor. If it be very hard, with a resinous fracture, it contains a large amount of shellac.

Lactarine. Sce Caseine.

Lactolin.—An acid lactate of potash, suggested by Boehringer as a substitute for lactic acid.

Lead Acetate.— $Pb(C_2H_sO_2)_2 + 3H_2O$ (Germ.: Bleizucker; fr.: Acétate de plomb, céruse de Mulhouse, saccharate de plomb, Sucre (sel) de Saturne). Synonyms: Normal or diacetate of lead, sugar of lead, salt of Saturn, "saccharum." Two kinds of sugar of lead are met with in the market—the brown and the white. The chief impurities which are likely to interfere with its use are acetates of iron and copper.

A coarser variety, known as brown acetate, is prepared by substituting distilled pyroligneous acid for the purer acid.

Lead acetate is only used for cotton. There it has been used for yellows and oranges in times long gone by. To-day a basic acetate with litharge, mordants cotton on simple immersion, and is used for eosines and similar colors.

Lead acetate has a sweet, pleasant taste, but is a virulent poison. Its antidote is Epsom Salts (which forms an insoluble lead sulphate), the whites of eggs, and emetics; avoid NaCl.

Lefebregummi.-Synonym for Brown Dextrines.

Leiocomme.—Synonym for Brown Dextrines.

Leiogomme.—Synonym for Brown Dextrines.

Lime, Caustic.—CaO (Germ.: Calciumoxyd, Gebrannter Kalk, Actzkalk, ungelöschter Kalk; fr.: Chaux vive (caustique, calcinée, anhydre). Synonym: Quick-lime.

Lime, Hydrate of.—CaH₂O₂ (Germ.: Calciumhydroxyd, Kalkhydrat, gelöschter Kalk; fr.: Chaux hydratée (éteinte), hydrate de chaux). Synonym: Slaked lime. Impurities are iron, aluminum, magnesium, and sand. Lime can generally be sufficiently tested by making a milk of lime, and observing the precipitate which is left; further, well-burned lime slakes and warms quickly, while badly prepared lime remains lumpy and gives off carbonic acid on addition of acids. From the slaking powers of limes they are characterized as "rich" or "fat," in contradistinction from "poor."

Litmus.—(Germ.: Lackmus; fr.: Tournesol.) Used as a test-paper. Alkalies turn it blue, while acids turn it red.

Logwood.—(Germ.: Blauholz, Campecheholz, Blutholz, Allerheiligenholz; fr.: Bois de campêche. Synonym: Campeachywood. Logwood is the wood of a South and Central American tree, *Hamatoxylum campechianum*. Chip-logwood sells for less than the logs themselves, because, after cutting up into chips, they increase in weight.

The analysis should embrace water, sp. gr., insolubles, alizarines, resins, glucose, hematin, and hematoxylin. Apt impurities are hemlock and chestnut extracts, molasses and glucose, farina, chlorides, sulphates, carbonates, insoluble residues, lime, etc. If chip-logwood is bought, test for anilines by throwing on to a beaker of water and noting how it dissolves. Use test-papers on the solution made. Look for moisture by weighing, drying, and reweighing; look for alkalis with litmus.

The comparative quality of samples of logwood may best be judged when they are dry. Those which exhibit the most stains and particles of a greenish bronze on the surface of the wood are the best. For greater accuracy, equal weights of the samples, say five grs. of each, may be taken and placed respectively in clear white glass phials or tubes of equal caliber and size. Upon each is poured 10 to 20 cc. of methylated spirit, and the phials are allowed to stand for an hour with occasional shaking.

For the comparative test dissolve 25 cc. of the sample to be tested in one liter of water, and in another liter of water the same quantity of the standard logwood; heat the two solutions to boiling, and then enter equal-sized pieces of cotton cloth; stir continually, and in about five minutes remove and squeeze carefully and evenly; do not wash, but throw them into a bath made up of 10 gr. bichromate of potash in a liter of water; stir well, and, after five minutes, heat so as to warm to about 30° to 35° Centigrade. Then wash well in cold water, dry, and compare. A logwood should be reported as follows:

Report No.
Laboratory No.
Hematine
Hematoxylin:
Ash (mineral):

Water:

Sp. Gr.: Beaumé: Date.

Their Mark or No.

Twaddle:

Insolubles:

Alkali:

Chestnut-extract:

Resins:

Alizarines:

Anilines:

Glucose:

A report of a logwood in this form places the management of the mill in a position to know whether the logwood is a proper logwood or not.

Lunar Caustic.-Nitrate of silver.

Madder.—(Germ.: Färberröthe, Krappwurzel, Röthe; fr.: Garance, racine de garance.) The root of Rubia tinctorum (peregrina, mungista), which formerly was grown in enormous quantities in the Levant and southern France. It is now entirely supplanted by alizarine. Its use was the dyeing of Turkey-red. The replacement of madder by products of coaltar is one of the most enduring monuments for the glory of chemists. It is rapidly assuming its true place as a museum euriosity.

Manganese Brown.—Cloth is saturated with MnCl₂ and then treated with NaOH, which must be free from Na₂CO₃—, otherwise manganous earbonate is produced, making the color irregular. One, Endler, passes the cloth impregnated with MnCl₂ through a bath containing 25 liters H₂O, 7 liters AmOH, and 500 gr. bichromate of potash. This produces an unstable chromate of manganese, which, on decomposing, allows the chromic acid to react on the manganous hydrate and change it to some higher state of oxidation.

Methyl-Orange.—(Germ.: Methylorange; fr.: Méthyl orange.) Dissolve one gram per liter. It is not affected by CO₂, and H₂S. Must not be used for organic acids, as the end reaction is indefinite. Can be used for the estimation of free acids in alum, ferrous-sulphate or chloride, zine-sulphate, copper-sulphate, or copper chloride.

Mordant.—(From Mordeo, I bite, Germ.: Beizmittel, Beize; fr.: Mordant.) Any substance used to bind the coloring-matter and the fiber. Vegetable fiber has not affinity for all dyes, and must be so treated as to combine with the dye to form a relatively insoluble compound.

Munjeet.—Synonyms: Mungeet, munjit, Indian madder. The root of Rubia munjista (cordifolia), a plant of the madder family. It comes in bunches of stalks nearly a yard in length, varying from the thickness of a finger downward. It does not contain much coloring-matter. It is used for making low qualities of garancine.

Myrabolans.—(Germ.: Myrabolanen; fr.: Myrabolanes.) A fruit produced in the East Indies, extensively consumed as a source of tannin, shape and size like a dried plum. The tannin is very variable in amount. Apt impurities are earth, sand, nux vomica, betel nuts, and a variety of seeds and berries, ground dividivi, old worthless sumac, and wild gall-nuts. The detection of these impurities is easily accomplished under the lens. A sample of the powder should be scattered on a white paper and a slight acquaintance with the different things used in sophistication will be revealed. Canaigre plant seems to be the better plant for tannin.

Nessler's Reagent.-See Ammonia.

Neuleim.-Different species of plant glue.

Oxycellulose.—This was discovered by Professor Witt. It can be made artificially by putting a piece of cotton into the solution of chemic and allowing it to stand for a little while, partially exposed to the air. Where the air strikes the cloth at the surface of the liquid, an oxidation product is formed, namely, oxycellulose. Its presence can be shown by boiling the piece of cloth in methylene blue. The oxycellulose has all the powers of a mordant for the aniline dye, and will show a deep coloration where the oxycellulose has been developed. Goods that are allowed to stay for any length of time in the chemic bin may have oxycellulose formed where the chemic solution has dried out.

Peachwood.—(Germ.: *Pfirsichholz;* fr.: *Bois de péche.*) This wood is one of the red woods similar in all its characters to Brazil-wood, although held to be poorer in coloring-matter.

Pearl-ash.—(Germ.: Perlasche; fr.: Perlasse.) Refined salt of potash (potassium carbonate, K.CO.).

Peroxide of Hydrogen.—H₂O₂ (Germ.: Wasserstoff superoxyd; fr.: Peroxyde (bioxyde) d'hydrogène). Used in such bleaching as feather-bleaching, but its cost has always limited its application.

Peroxide of Sodium.-Na2O2 (Germ.: Natriumperoxyd; fr.:

Peroxyde de soude). A bleaching agent manufactured on a large scale, and successfully used on the bleaching of fibers, particularly of mixed goods. Dr. Haen has proposed that the peroxide be used in conjunction with Epsom salts.

Phenacetoline.—For KOH or NaOH in carbonates.

Phenol-Phthaleine.—For titrating oxalic, acetic, citric. or tartaric acids. Five gr. per liter is proper strength. Crimson when alkaline, colorless when neutral or acid.

Pink Salts.—SnCl₄,2AmCl (Germ.: Pinksalz, Ammonium-Zinnehlorid; fr.: Chloride d'étain ammoniacal). A name given to the double chloride of tin and ammonia. It was formerly employed instead of the other salts of tin, in the wood-pinks; it is now very seldom met with in commerce.

Plant-Glue, different species of:

Alligin, Hydrofugicolle,

Arabil, Japanleim, Brilliantkleister, Neuleim, Gloriagummi, Traganthine,

Gummi Germanicum, Universalleim.

Gummi Saxonicum.

Plaster of Paris.—CaSO₄ (Germ.: Gebraunter Gips; fr.: Plâtre fin). Synonyms: Sulphate of lime, or white mineral.

Potash.—K₂O (Germ.: Kaliumoxyd. Kali; fr.: Potasse. Synonym: Potassa. It should dissolve entirely in pure water, leaving no sediment or residue. If dissolved in distilled water, acidified with nitric acid, the solution, on the addition of nitrate of baryta, should not give a precipitate. If one appears, it denotes the presence of sulphuric acid, present as sulphate of potash.

Potash, Chlorate of.—KClO₃ (Germ.: Kaliumchlorat, Chlorsaures Kali; fr.: Chlorate de potasse). The solution should yield no precipitate with a dilute solution of the nitrate of silver.

Potash, Chromate of.— $K_2\mathrm{CrO}_4$ (Germ.: Kaliumchromat, Chromsaures Kali; fr.: Chromate de potasse). Synonyms: Yellow chrome or neutral chrome. The impurities in yellow chrome are carbonate of potash and sulphate of potash. The latter is sometimes present to the extent of 56 per cent.

Potassium, Bichromate of.—K₂Cr₂O₇ (Germ.: Doppettchrom-saures Kalium, Kaliumbichromat; fr.: Bichromate de potasse). Apt impurity, sulphates.

Potassium, Prussiate of.—KCy or K(CN) (Germ.: Cyankalium, blausaures Kali; fr.: Prussiate de potasse, cyanure de potassium). Synonyms: Prussiate of potassa (potash), cyanide of potash. Very poisonous. Antidotes: Emetics; cautiously inhaled H₃N.

Potash, Yellow Prussiate of.—K₄FeCy₆ (Germ.: Ferrocyankalium, Kaliumeisencyanür, Gelbes Blutlaugensalz; fr.: Prussiate jaune de potasse, cyanoferrure de potasse, ferrocyanure de potasse, lessive du sang). Synonyms: Yellow prussiate, potassium ferrocyanide, ferrocyanuret of potash, blood-salt. The chief contamination to which it is liable is sulphate of potash. It is used for making "Prussian Blue."

Potash, Red Prussiate of.—K₃FeCy₆ (Germ.: Ferricyankalium, Kaliumeisencyanid, Rothes Blutlaugensalz; fr.: Prussiate rouge de potasse, cyanoferride de potasse, ferricyanure de potasse).

Quercitron Bark.—(Germ.: Quercitron Rinde; fr.: Quercitron.) The bark of Querens tinctoria (nigra). Synonym: Bark. It was introduced into England at the close of the last century, and soon came into use as being cheaper and stronger than the yellow coloring-matters then known in the trade. It dyes up good yellows upon wool and cotton—on the first with a tin mordant, and upon the second with an alumina mordant. Quercitron is ballasted with NaCl, Na₂CO₃, etc.

Red Liquor.—(Germ.: Essignaure Thonerde, Aluminiumacetat; fr.: Acétate d'alumine.) Synonyms: Acetate of alumina, red mordant, pyrolignite of alumina. An aluminium sulphoacetate appears to act satisfactorily. It is prepared by mixing, first, 453 pounds ammonia alum (383 pounds aluminium sulphate), 379 pounds lead acetate, 1,132 pounds water; or, second, 453 pounds alum (or 333 pounds aluminium sulphate) and 158 pounds acetate of lime. The mixture is agitated, settled, and the clear liquid decanted.

Red Liquor usually contains from 3 to 5 per cent, alumina $(\mathrm{Al}_2\mathrm{O}_3)$ and 6 to 10 per cent, acetic acid; its density varies from 1.085 to 1.120.

Redwood. See Brazil-wood.

Salt (common).—NaCl (Germ.: Kochsalz, Chlornatrium; fr.: Sel). Apt impurities are CaSO₄, MgCl₂, NaSO₄, and insolubles.

Santal-Wood .- (Germ.: Sandelholz, Rotes Santelholz, Calia-

turholz; fr.: Bois du santal, bois de santal rouge.) Synonyms: Sandal-wood, Saunders-wood, and Red-sanders, Pterocarpus santalinus (indicus).

Soap.—(Germ.: Seife; fr.: Savon.) Additions are glycerine, derived from the oil or fat employed, water, alkaline sulphates, chlorides, and carbonates from the impurities in the potash or soda employed, rosin, silicate of soda, Cornwall-clay, ground flints, potter's slip, Fuller's earth, gelatine, and other kinds of nitrogenous animal matter, dissolved in soda; farina, dextrine, and a variety of other substances.

Rosin is a common ingredient in yellow soaps. In all soaps intended for the use of the dyer or printer, it is an objectionable impurity, imparting to the goods an unpleasant elamminess or stickiness. It may be easily detected. Dissolve some of the soap in as little boiling water as possible. Add enough hydrochloric, or dilute sulphuric, acid to combine with the alkali, and liberate the fatty matters. Then cover the vessel in which the whole is contained with a lid or a plate of glass, and set it aside for a few minutes. On uncovering it, rosin, if present, may be easily detected in the cake of fatty substances by its smell and taste, and by the peculiar stickiness which it imparts to oils and fats with which it is mixed.

Silicate of soda as a constituent of soap has given rise to a great variety of opinions. Some condemn it as a mere adulteration and even deny that it has any detergent powers.

Insoluble silica and alumina, in the various shapes of Cornish-clay, powdered pumice, ground flints, potter's slip, and Fuller's earth, can be and often are added to soaps to a considerable extent.

Animal matter other than fat, such as tendons, intestines, ground bones, waste glue, etc., can be regarded in no other light than a filthy adulteration. The detection of such impurities is not difficult. A little of the soap is dissolved in hot water, and decomposed by the addition of a little hydrochloric acid. The spurious animal matters will be entangled in the cake of fat which separates out, and may be easily recognized.

Farina and dextrine are less formidable. They add, of course, nothing to the value of the soap, but they have no positively injurious action. To detect them, dissolve the soap in strong alcohol. Farina and gum remain undissolved, and

may be separated from the other articles by filtering. Farina is then easily recognized by adding a drop or two of the tincture of iodine, which, if farina be present, will give to the mass a deep blue-black color. Dextrine (British gum) may be dissolved out from any mineral impurities present in a little hot water, and can be readily recognized.

The following oils and fats should be carefully avoided: Train oil, cod-fish oil, linseed oil, kitchen refuse, fat collected by bone boilers, grease separated out from accumulated soaplyes. All these give evil-smelling, clammy soaps.

Soap Powders.—Washing powders, scouring salts, washing sugars, soap ashes, extracts of soap, extract of Fuller's earth, saponaceous, etc., etc. These are bodies which have been very largely used, not only for domestic, but also for manufacturing purposes.

Soda Ash.—Na₂CO₂ (Germ.: Soda, kohlensaures Natron; fr.: Soude, carbonate de soude). Synonyms: Soda carbonate, ash, alkali, or soda. Apt impurities: Substances insoluble in water, chlorides, phosphates, sulphates, lime, iron, sulphides, sulphites and hyposulphites, and caustic soda.

Perhaps the most harmful impurity is caustic soda, which is very injurious to wool. Test for this by precipitating the carbonate with barium chloride, filter, and add phenol-phthalein; a pink color is developed if caustic soda was present.

The sulphite and hyposulphite are tested for by adding some dilute sulphuric acid, to which a little bichromate of potash has been added. If a green color is produced the solution is proved to have held the sulphite or hyposulphite.

The sulphide of sodium is tested for by the addition of sulphuric acid, when the odor of H₂S is evolved; if a portion is dissolved in water and then filtered, produces a black precipitate (PbS), when a solution of acetate of lead is poured into it.

The strength is found by titration simultaneously with the standard. Both samples must be dried equally, dissolved in distilled water in exactly equal weights, and methyl-orange used as the indicator.

Soda ash is usually sold on a basis of 48 per cent.; for every per cent, over 48 per cent., 1/48 is added to the bill. This full claim is very rarely sustained by the analysis. Mills should buy on sample and then send the sample by which they purchased to the laboratory, and also a small sample taken from

the lot shipped them, and then the bill should be paid in accordance with the results of the analysis.

The normal solution of soda is 53 gr. per liter.

Soda Crystals.—Na₂CO₃ + 10H₂O (Germ.: Krystallisirtes kohlensaures Natron, Sodasalz; fr.: eristaux de soude, sel de soude). Synonyms: Carbonate of soda crystals, Scotch soda, Newcastle soda, crystallized carbonate of soda, washing soda, sal soda. Contains 63 per cent. of its weight of water.

Soda, Sesquicarbonate of.—Na₂CO₃, 2NaHCO₃ (Germ.: Anderthalbfach-kohlensaures Natron; fr.: Sesquicarbonate de soude). The principal constituent in natural "Trona" ("Urao") Soda. As it is anhydrous, it contains nearly 49 per cent. Na₂O, which is equivalent to 83 per cent. Na₂CO₃.

Sodium, Arseniate of.—Impurities: Salt and phosphates. See Arseniates.

Sodium, Bisulphite of.—NaHSO₃ (Germ.: Zweifach-schwe-tligsaures Natron; fr.: Bisulfite de soude). See Antichlor.

Sodium Hyposulphite (thiosulphate) of.—Na₂S₂O₃ + 5H₂O (Germ.: *Unterschwestigsaures Natron*; fr.: *Hyposulfite de soude*). Synonym: Green mordant.

Sodium, Hydrate of.—NaHO (Germ.: Naturiumhydroxyd, Aetznatron; fr.: Soude caustique à la chaux). Synonym: Caustic soda. Impurities can be NaCl, Na₂SO₄, alumina, silica, carbonate, and iron. Titrate with methyl-orange.

Sodium, Silicate of.— Na_2SiO_3 (Germ.: Kiesclsaures Natron, Wassergluss; fr.: Silicate de soude). Synonyms: Soluble glass, waterglass, flint-liquor. A compound formed by melting together sand with carbonate of soda ($SiO_2 + Na_2CO_3 = Na_2SiO_3 + CO_2$). It is used as an addition to soaps; as a sizing and as a mordant. Combines the properties of soap and caustic alkali, and is well adapted to some operations where the sodaash is not strong enough and where the alkali is too keen.

Sodium, Stannate of.—Na₂SnO₃ + 3H₂O (crystals) (Germ.: Zinnsaures Natron, Grundirsalz, Präparirsalz; fr.: Stannate de soude). Synonym: Preparing salt. To ascertain its value, a portion of the sample should be dissolved in a small quantity of water, slightly acidulated with hydrochloric acid, and some strips of clean, pure metallic zinc placed in the solution. The tin will be thrown down in a spongy state.

The impurities likely to be met with are common salt—an intentional fraud—and an excess of alkali, caustic or carbon-

ated, resulting from bad workmanship, of nitrate of sodium, sulphate of magnesium, and iron. Certain additions are also occasionally made to the stannate, not with fraudulent design, but avowedly for the purpose of improving its quality. These are the tungstate of soda, the antimoniate of soda, and the arsenite and arseniate of soda. Sodium stannate should dissolve clear, but not too clear, in which case suspect free muriatic acid; its value depends upon the amount of tin it contains.

Method of Analysis.-The sample must first be ground in a mortar, and then anywhere from two to five gr. of the stannate is weighed out and ground again in a mortar with the water. The mixture is allowed to stand for about half an hour, then it is poured on to a weighed filter-paper. It is washed with warm water several times until the wash-water going through the filter gives no precipitate when acidified. The filtrate contains all the soluble stannate of soda. The filtrate is heated with dilute sulphuric acid, running the acid in from a burette: the acid solution clouds up, and must be allowed to settle, after which the clear liquid is poured off and filtered; the precipitate is washed once with warm water and then the whole is washed on to the filter-paper by means of a wash-bottle. It is washed three or four times on the filter-paper, dried, placed in a porcelain crucible, heated to red heat, cooled, moistened with nitric acid, heated again to a red heat, and weighed.

Sodium, Sulphate of .- See Glauber's Salts.

Starches. Analyses and Tests.—The relative tenacity of starches can be determined by weighing out equal parts of each sample, mixing with equal volumes of water, boiling briskly for three minutes, constantly stirring, and then pouring into a conical test-glass and allowing to cool. Then carefully turn the contents of each glass out, and note time it takes before a "land-slide" occurs. The starch which stands longest is best.

Starch is sometimes deteriorated by the presence of nitrogenous matter. To detect this, dissolve one part of mercury in two parts of nitric acid of 50° Twaddle at a gentle heat. When dissolved, boil the liquid for a few minutes. If the sample is colored reddish when boiled with this reagent, nitrogenous matter is present. In the laboratory the following determinations are made: Ash, viscosity, moisture (allowing

15 per cent.), Fehling's solution test, grit, taste; if there is considerable ash, the starch is tested for sulphates, chlorides, mineral-white, gypsum, and china clay.

Pure wheat-flour is almost entirely dissolved in a strong solution of potash containing 12 per cent. of the alkali; but mineral substances used for the purpose of adulteration remain undissolved. Under the microscope the granules are characteristic and very well defined. In some cases the microscopic examination is aided by an application of a solution of potash. Thus we may readily detect the mixture of wheat-flour with either potato-starch or meal of peas or beans by the addition of a little water to a small quantity of flour, then by adding a few drops of a solution of potash, when the granules of the potato-starch will immediately swell up and acquire three or four times their natural size, while those of the wheat-starch are scarcely affected; if adulterated with pea- or bean-meal, the hexagonal tissue of the seed is at the same time rendered very obvious under the microscope.

The reaction of starch ought to be neutral; if it shows alkaline reaction it should be rejected, as it is in that case apt to cause trouble in finishing. The formation of a gelatinous paste is accelerated by adding caustic potash, and for that reason some dealers add this impurity.

The usual method for the determination of starch is to invert by the action of dilute acid, nitric acid, and then determine the dextrose produced by Fehling's solution. One hundred parts of the dextrose are taken as equivalent to ninety of starch.

The proportions for the iodine test are: One gram starch stirred to a cream with a little cold water; then add 60 cc. hot water, and the whole is boiled; of this mixture, three drops are placed in 10 cc. water. and the iodine colors it as soon as it is set free.

Potato-starch produces the thickest paste, weight for weight, because the grains are exfoliated by high heating. Rice is less adhesive, and cannot be used to fix large proportions of earthy matter, clay, or minerals in the fabric.

Test for Flour in Starch.—Stir to a paste with strong nitric acid, and if the sample be starch it will remain white, but if there be any flour in the sample the color will alter to a

strong yellow. The sp. gr. test often gives interesting information. It can be made by boiling a certain weight of a sample in sufficient water to prevent it from solidifying when it cools; the sp. gr. of the cool solution gives a comparative test of oftentimes considerable value.

The adulteration of mineral matter is best detected by shaking the sample in a test-tube with choloroform. The starch floats on the chloroform and the mineral substances go to the bottom.

In German the different starches are known as follows: Potato is Fecule; cor is Mais; wheat-starch is Weizen.

Sumac.—(Germ. Sumach, Schmack; fr.: Sumac.) Synonyms: Sumach or shumac. Leaves and smaller branches of a shrub, Rhus coriaria (Cotinus). That which comes from Sicily is the most esteemed, and brings the highest price, but several other countries produce a usable article. Has a smell reminding of tea, or sometimes of new hay. Its quality can be judged of by its color to a considerable extent; it should be bright and clear; some samples are dull and have a faded brown look. Examination should embrace the ash, water content. insolubles, glucose, dyeings on unmordanted cotton and wool. Sumac has the same chemical properties as galls, containing the same acids, tannic and gallic; but, in addition, it has a certain amount of yellow coloring-matter, which, though nearly worthless in itself, modifies its effects upon mordanted cloth.

The chief consumption of sumae is probably in cotton-dyeing, where it is the preliminary treatment for nearly all the fancy shades to steep the cotton for some hours in decoction of sumae. The astringent matter of the sumae is thus firmly combined with the cotton, which can now be easily mordanted with either tin or alumina, which forms the basis of the colors. Sumae liquors have a strong tendency to become acid, which must be guarded against in those cases where an iron-alumina mordant is concerned, since the acidity is sometimes strong enough to dissolve out weak iron mordants.

The laboratory method for reporting sumac is as follows: Report No. Date.

Laboratory No. . Their Mark or No.

Tannic acid:

Ash (mineral):

Water:

Specific Gravity:

Beaumé:

Twaddle:

Insolubles:

Resins:

Anilines:

Alizarines:

Glucose Test:

Tartar-Emetic.— $K(SbO)C_4ll_4O_0 + \frac{1}{2}H_2O$ (Germ.: Brechweinstein, Spiessglanzweinstein; fr.: Tartar émétique). Synonyms: Potassium-antimony-tartrate, tartarus stibiatus. Mixtures of tartar-emetic for mordanting are sold containing as much as 33 to 59 per cent. zinc sulphate at a lower price, under the names tartar-emetic powder, tartar-emetic substitute, antimony mordant, etc. It is known that zinc acetate may partly replace the tartar-emetic with advantage, but the sulphate appears to be a simple adulterant.

Several other compounds of antimony have been proposed instead of tartar-emetic for mordants. E. Jacquet advises the use of a mixture of basic antimony oxalate with twice its weight of ammonium oxalate. Nölting recommends the double oxalate of potash or of ammonia. The use of the fluoride (which is not precipitated with the excess of water) and the double fluorides of antimony and the alkalies has been patented by S. M'Lean. C. Watson, Jr., has recently patented a process for using tri-chloride with sufficient common salt to prevent the precipitation of the oxychloride.

Test.—Take 1 gr. of the sample to be analyzed, add starch solution, and ½ to ¼ gr. Na₂CO₃, 10H₂O, then let the centi-normal iodine solution run in; shake till the blue disappears, and add more until the blue remains permanent, which will occur when all antimonoxide is converted to antimonic acid. Multiply the cc. of iodine solution used by 1.66, and the per cent. of tartar-emetic is found; 0.72 is per cent. of antimony oxide.

A double salt of antimony-fluoride and ammonium sulphate, SbF₃.(NH₄)₂SO₄, is patented and sold by Haën, of Hanover, Germany, and is called by him "Antimony Salt." It forms white oblique crystals, and dissolves very easily in water. The solution possesses a strong acid reaction, and attacks glass and metals like other fluorides. It contains 47 per cent.

Sb₂O₃. Nine parts of it are equal to 10 parts of tartar-emetic. Tartar-emetic has 43 per cent. Sb₂O₃. Antimony oxalate contains 23 to 27 per cent. oxide of antimony.

Tartar, Crude. See Argols.

Tartar, Cream of.—When used as a mordant cream of tartar should be employed to the extent of 2 per cent., then bichromate of potash, 3 per cent.; this should be boiled one hour, cooled, and washed.

Tin Mordants; Stannous Chloride.— $\operatorname{SnCl}_2 + 2H_2O$. (Germ.: Zinnehlorür, Zinnsalz; fr.: Sel d'étain). Synonyms: Protochloride of tin, "tin crystals," or "tin-salt." Apt impurities are sulphate of zinc, sulphate of magnesia, and chloride of zinc. When sold in solution it is known as the single or double muriate of tin, according to the specific gravity 1.3 or 1.6; 60° or 120° Twaddle.

Stannous chloride and certain dyewood extracts form "Carmines," "Persian Berry Carmines," "Cochineal Carmine," etc., occasionally used. In Turkey-red the tin crystals are used to brighten the color. In the case of wool the tin crystals are used with cream of tartar and other assistants. In silk-dyeing it is used principally in conjunction with catechu in producing heavily weighted blacks.

"Tin Spirits" is a name given to solutions of tin, in the manufacture of which other acids besides HCl are used, e.g., HNO₃ and H₂SO₄. They are variable in composition, and their value must be determined especially in each case. One of them is "Scarlet Finishing Spirit," a muriate of tin containing a certain proportion of oxalic acid and sometimes tartaric.

Toluidine.—Orthotoluidine (formerly known as pseudotoluidine) is an oily liquid, boiling at 386° F. It gives a violet color with solution of chloride of lime and muriatic acid, and a blue with a mixture of sulphuric and nitric acid. Metatoluidine, which is less often met with, is also a liquid boiling at the same temperature. These toluidines give each different results in color-making.

Traganthine.—A different species of plant glue.

Turmeric.—(Germ.: Gelbwurzel; fr.: Safran des Indes, souchet des Indes.) Synonyms: Curcuma, Terra merita. A yellowish coloring-matter, the root of Curcuma longa rotunda, growing in the East Indies, and now chiefly cultivated in Bengal. It is mostly used in silk-printing and dyeing. Is used

also for dyeing the cotton employed for making ham-bags. The color is not at all fast. The only adulteration to which turmeric is liable in commerce is common salt, which is sometimes added in quantity to the roots while going through the mill.

Ultramarine.—(Germ.: Ultramarin; fr.: Bleu d'outremer.) Synonyms: Lazurstein, lapis lazuli, silicate of alumina and soda. Impurities might be coal-tar colors, Turnbull's blue, gypsum, and moisture. The most satisfactory test is a comparative test showing the relative resistance of a sample against acid and alum.

Universalleim.—Different species of plant glue.

Weld.—(Germ.: Wau; fr.: Gaude.) Synonym: Wold. The yellow dyeing-color employed before the introduction of quercitron. It is the herb Reseda ruteola, and sold in the sheaf, like straw; the whole of the plant, except the roots, were employed in dyeing, but the greater part of the color resides in the seeds and upper extremity.

Whitewash.—(Germ.: Kalkbrühe, Weisse; fr.: Lait de chaux.) The best proportions for whitewash are two pails of water, ¾ pound quick-lime, 1½ pounds salt, and 1 pound of dextrine.

Woad.—(Germ.: Waid, Pastel; fr.: Pastel, vouède, guède.) This dyeing matter, which was employed from the most ancient times, is now nearly unknown in this country. It is yet eultivated in some parts of Europe (Isatis tinctoria, Nerium tinetorium), where it goes under the name of pastel. The coloring-matter it contains is chemically and practically the same as indigo; it is still used in setting the indigo vats for dyeing woolen, but always in conjunction with indigo. It appears that the woad plant, as sold to the indigo dyer, readily enters into fermentation, and in that state is useful in deoxidizing or reducing the indigo to the soluble condition; but it contains very little coloring-matter itself, so that it was hardly possible to dye a deep blue with it.

Xanthopyll. See Chlorophyll.

Zinc Acetate.— $Zn(C_2H_2O_2)_2$ (Germ.: Essigsaures Zink; fr.: Acetate de zine). Recommended as a substitute for tartaremetic in fixing tannin on cotton. It is also used in wooldyeing with malachite-green.

Zinc Dust .- (Germ .: Zinkstaub; fr.: Poussière de zinc.)

The ZnO₂H₂ always present in specimens of zinc-dust may be removed by digesting with NH₄Cl-water, and then with NH₂-water and drying.

Zinc Chloride.—ZnCl₂ (Germ.: Chlorzink; fr.: Chlorure de zinc). Synonyms: Muriate of zinc, butter of zinc. The use of zinc chloride is steadily decreasing. It is antiseptic and deliquescent.

Zinc Sulphate.—Sulphate of zinc.

PART III.—NEW BOOKS.

"Matières Colorantes Artificielles" is a two-volume work of 1,645 pages, by Léon Lefèvre, and should be in the hands of every color-chemist. In many ways it is an advance on Schultz's "Chemistry of the Coal-Tar Colors." The last French book that went into the subject with anything like thoroughness was the three-volume work of Depierre. The book is fully illustrated with samples dyed and printed on cotton, wool, and silk. It is published by the house of G. Masson, of Paris.

The third edition of Schultz's "Tabellarische Uebersicht der Künstlichen Organischen Farbstoffe" appears from the press of R. Gärtner, Berlin. It is impossible for anyone who pretends to keep abreast with the times in the matter of the coal-tar colors to get along without this book. Each dye is given with its trade name, its scientific name, and its constitution, the literature concerning the dye, the properties, reactions, and application. It is altogether the best and most usable book which has ever appeared in the technology of dyes and dyeing.

"Dictionary of Coal-Tar Colors," by J. H. Hurst. This is a translation of Schultz & Julius. It is a great misfortune for a colorist not to have a practical command of German and French, and particularly the German. Almost all the information is published in German, sometimes French in the first instance. For those who have not a practical control of German the Hurst "Dictionary" is a very valuable book to have. The edition to hand is the second edition.

"Färbereichemische Untersuchungen," by Dr. Paul Heermann, Berlin; Julius Springer, 1898; pp. 138. Priee, 4 marks. This is a well bound and useful book to the color chemist. In pocketable shape it places in the hands of the chemist the data desired and of frequent application. It is more and more becoming evident that our manufacturers are being driven to

the testing and examination of the dyes, chemicals, and drugs which they use. This book furnishes many short and direct methods. The chapter on the fibers is excellent. The legal test for cotton in woolens is given.

"Mikrochemische Analyse organischer Verbindungen," by H. Behrens. Published by L. Voss, of Hamburg. Price, 4.50 marks.

"Yarns and Textile Fabrics," by Dr. J. Korzfeld, translated by Charles Salter, London. Scott, Greenwood & Co., 23 Ludgate Hill, London, E. C.; pp. xiv., 207. This excellent book, upon a subject in which all manufacturers are necessarily interested, comes to band in excellent shape. On pages 5 and 6 it says: "In the work are collected all the tests, both of a physical and chemical nature, that can, for technical purposes, be performed on yarn or cloth, so that not only the commercial and textile chemist who has frequently to reply to questions of this kind, but also (and principally) the practical manufacturer of textiles and his subordinates, officials, overlookers, the spinner, the weaver, the dyer, and the finisher are catered for."

"Chemische Technologie der Azofarbstoffe," by Dr. Carl Bülow. Leipzig: Otto Wigand, 1897. The first part, 123 pages, gives a highly scientific treatment of the azo-dyes. The author arranges the azo-dyes into an order and classification which seems to be a step in advance of what has hitherto been habit and custom. As the work is of a highly scientific nature, it is a little beyond the limits of a publication of this nature to give any very extensive review of the work. When the whole work is completed it will again be brought to attention, and will be commended for all those chemists who wish to become perfectly familiar with this, perhaps the most extensive group of all the coal-tar colors.

"Die Appreturmittel und ihre Verwendung," second edition, by Friedrich Polleyn. Vienna, Pesth, and Leipzig: A. Hartleben. This is a brochure of 250 pages on the general subject of the sizing of textiles. It is of a pocket size, and comes along in a set of technological books which have been issued for a long time by the press of A. Hartleben. The formulæ are very well grouped into classes. The more modern sizing methods have received generous notice, and the methods for making fire-proof and water-proof finishes are

stated quite fully. At the end of the volume suggestions for the analysis of sizings are given.

"Sizing Ingredients, Size Mixing, and Sizing," by Hugh. Monie, Jr. Manchester, England: Abel Heywood & Son, 56 Oldham Street. 63 pages. Illustrated. Price, \$1.00. The author of this work is well known as the author of "Cotton and the Cotton Fiber," etc., and editor of the *Indian Textile Journal*, Bombay.

"Modern Bleaching and Finishing," by John Heywood, Manchester, England.

"Mechanische Weberei-Praxis," by Franz Donat.—This book of 144 pages is divided into three parts, the first being the mechanical idea of weaving in general; the second part discusses the different looms; and the third section is a specially interesting one, giving the numbering and certain details concerning not only cotton-yarn, but linen, hemp, jute, China-grass, woolen yarns, etc. There is an appendix, giving the tables of calculation and interesting details concerning the mechanism of yarn.

"Bindungs-Lexikon für Schaftweberei."—This book comes bound and, together with the foregoing, should be in the libraries of our different textile schools. The Lexikon contains 160 pages, printed in red and black, giving valuable information in concentrated form for the weaver. The book is handsomely bound in imitation leather, and the presswork is very creditable.

"The Blue Book," 1898 edition, appears in its usual attractive and complete form. This publication is so well known to the trade that it needs nothing more but a notice of its desirability in the columns of the Year-Book.

"Abhandlung über das Färben mit Blauholz-Extract."— This is a publication issued by the Leipziger Farbwerke Paul Gulden Company. It is a small book, but seems to us to be very valuable, giving, as it does, the dyed samples and illustrating the advanced methods of testing for hematine and hematoxylin. It is worth while sending for this book.

"Technical Education."—The Society for the Encouragement of Arts, Manufactures, and Commerce, John Street, Adelphi, London, W. C., England, has published a "Report of the Proceedings of the late International Congress on Technical Education," held in London last June. Among the papers re-

ported may be mentioned "Relations Existing Between the Teaching of Pure Chemistry and Applied Chemistry."

"The Printing of Textile Fabrics: A Practical Manual on the Printing of Cotton, Woolen, Silk, and Half-Silk Fabries," by C. F. Seymour Rothwell, Lecturer on Calico and Delaine Printing at the Municipal Technical School, Manchester, England. Charles Griffin & Co., Ltd., Exeter Street, Strand, London, 1897, and J. B. Lippincott Company, Philadelphia, l'a. Price, \$6.50. It is said that this book is intended to form a companion volume to Knecht, Rawson, and Loewenthal's Manual of Dyeing. There are over 300 pages, followed by 20 pages of printed and dyed patterns. The following subjects are treated: Introduction. Methods of Textile Printing. Styles. I. Machinery used in Textile Printing. II. Thickeners and Mordants. III. Printing of Cotton Goods. a. Bleaching of Cotton Piece Goods for Printing. b. The Steam Style. c. Colors Produced Directly on the Fiber. d. Dyed Styles. c. Padding Style. f. Resist and Discharge Styles. g. Printing of Compound Colorings. IV. Printing of Woolen Goods. V. Printing of Silk Goods. VI. Practical Recipes for Printing. Appendix of Useful Tables.

"Elements of Chemistry," by Rufus P. Williams. New York: Ginn & Co. An excellent work of 400 pages. Mr. Williams says that he wishes the learner "to think scientifically."

"Theoretical Chemistry," by Ira Remsen. New fifth revised edition. Price, \$2. Philadelphia, Pa.: Lea Brothers & Co.

"Recent Progress in the Industries of Dyeing and Calico Printing: Being a Supplement to The Printing of Cotton Fabrics and Dyeing, by Antonio Sansone. Manchester, England: Abel Heywood & Son. A chapter in this book is devoted to the use of colors produced upon the fiber. This important question is treated in detail, and a great many interesting facts are given. One chapter is given to cop-dyeing.

"The Anthracene Dyestuffs." By R. Gnehm. Braunschweig: Vieweg & Son.

"Anleitung zur Qualitativen Chemischen Analyse," by R. Fresenius. Sixteenth edition. Braunschweig: Fr. Vieweg & Son. Price, 12 marks.

"Handbuch der Soda-Industrie und ihrer Nebenzweige," by G. Lunge. Zweiter Band. "Sulfat, Salzsäure, Leblancverfahren." Second edition. Braunschweig: Fr. Vieweg & Son. Price, 30 marks.

"Lehrbuch der Farbenchemie," by G. v. Georgievics. Leipzig: Fr. Deutieke.

"Die Künstlich erzeugten Organischen Farbstoffe, by Rich. Meyer. Braunschweig: Fr. Vieweg & Son.

"Chemie der Organischen Farbstoffe," by R. Nietzki. Berlin: Julius Springer. Second edition. Price, 8 marks.

"Fortschritt der Theerfarbenfabrikation und verwandter Industriezweige," by P. Friedländer. Berlin: Julius Springer. Price, 40 marks.

"Die Industrie der Stärke und der Stärkefabrikate in den Vereinigten Staaten von Amerika und ihr Einfluss auf den Englischen Markt," by O. Saare. Berlin: Julius Springer. Price, 3 marks.

"Technical Lexicon in Four Languages." Vol. I. German, Italian, French, and English. Vol. II. Italian, German, French, and English. Vol. III. French, Italian, German, and English. Vol. IV. English, Italian, German, and French. By Edward Webber. Berlin: Press of Julius Springer. Price, per volume, 75 cents. Each volume contains 8,000 technical words and expressions. The books are not large, and make no pretense to universality, but they are most useful to those who are not trained in the trade phrases and expressions in the technique.

"Year-Book of the Oil, Paint and Drug Reporter." This is a brochure of 83 pages, and is filled to overflowing with interesting statistics and particulars relative to the special field covered by the Oil, Paint and Drug Reporter. The tariff acts of 1890, 1894, and 1897 are given in tabular form, in so far as tariff acts refer to oils, paints, chemicals, and dyestuffs.

"Lubricating Oils, Fats, and Greases: Their Origin, Preparation, Properties, Uses, and Analysis," by George H. Hurst. Ludgate Hill, London: Scott, Greenwood & Co., 1896. Price, 10s. 6d. An octavo volume containing frontispiece, preface, table of contents, and subject-matter filling 301 pages, four pages of tables, and an alphabetical index. The work is illustrated with 65 wood-cut illustrations. The text is divided into eight chapters, entitled as follows: 1. Introductory, Oils and Fats, Fatty Oils, Hydrocarbon Oils, Uses of Oils. 2. Hydrocarbon Oils. 3. Seotch Shale Oils. 4. Petroleum. 5. Vegetable

and Animal Oils. 6. Testing and Adulteration of Oils. 7. Lubricating Greases. 8. Lubrication.

"Cotton: Its Uses, Varieties, Fiber Structure, Cultivation, and Preparation for the Market, and as an Article of Commerce; also the Manufacture of Cotton-seed Oil, Cotton-seed Meal and Fertilizers, with Especial Reference to Cotton Growing, Ginning, and Oil Pressing in the United States," by C. P. Brooks. New York: Spon & Chamberlain, 12 Cortlandt Street. This is as attractive a book on a technical subject as it has ever been our pleasure to handle. It is more than ordinarily interesting in its get-up and pictures, and is, in addition, beautifully edited. It is handsomely bound, of almost four hundred pages, and is most profusely illustrated.

"Lehrbuch der Chemischen Technologie der Gespinstfasern," Part II., by Dr. George von Georgievics. Leipzig and Vienna: Franz Deuticke. The second part of this book contains 354 pages, treating particularly of fibers, bleaching, dyeing, printing, and sizing. The book is an excellent one on all of the operations pursued in bleaching, dyeing, and printing establishments. The mordants are discussed in detail, the dyeing methods, tests of the dyes, and the dyed fiber—altogether it is one of the books which it is advantageous to have.

"Dictionary." The Société Industrielle de Mulhouse having charged itself with the publication of the dictionary of M. Jules Garçon, announces that the book will appear shortly. It will give a methodical arrangement of the different departments concerning textile industry, and in alphabetical order will mention all the researches made in the scientific journals, and the reviews on dyeing and calico-printing. The value of such a work can scarcely be over-estimated.

"Dictionnaire des Altérations et Falsifications des Substances alimentaires, medicamenteuses et commerciales, avec l'indication des moyens de les reconnaître," by A. Chevallier and E. Baudrimont. Seventh edition, by L. Herbert. Vol. II. Paris, 1897. Svo, with figures.

"Sur le Tannin de la Galle d'Alep et de la Galle de Chine." by E. Manceau. Epernay, 1896. Pp. iv., 147. Price, 6 marks.

"The Diamine Colors." The firm of William J. Matheson & Co., Ltd., has published a book of diamine colors, in the usual generous and handsome style in which their publications appear. It is exclusively occupied with the diamine colors, is handsomely bound, and consists of about 250 pages.

The firm of Victor Koechl & Co. has published a handsome book on the coal-tar colors. The publication follows somewhat the line of the classical Schultz & Julius. The range of colors is very complete, and leaves little to be desired.

"Bleaching and Finishing Machinery for Textile Fabrics."

—H. W. Butterworth & Sons Company, Philadelphia, Pa., have issued a trade catalogue of 28 pages, relating to their machinery for bleaching, printing, dyeing, drying, and finishing of textile fabrics. The catalogue contains many illustrations of the machines of this concern.

"A Descriptive Catalogue of Useful Fiber Plants in the World," by Charles Richards Dodge. Washington: Department of Agriculture. 362 pages, 13 plates, and 103 text illustration. This is a descriptive catalogue of the useful fiber plants of the world, including structural and economic classifications of fibers. Mr. Dodge is recognized as an authority in this country upon fiber plants.

"The Cotton-Plant."—The United States Department of Agriculture issues in book form several of the valuable bulletins of their Experiment Stations. The price is 60 cents, which covers the cost of printing and binding.

The State Geologist of Georgia has compiled a report of the water-powers of Georgia. There are many unused waterpowers in the Piedmont district of the South Atlantic States. Those interested in the question of mills in that section should obtain this report. The water-powers of Georgia are confined to what is termed the Crystalline Belt of the State.

"Some Notes in Chemical Jurisprudence; An Abstract of Cases of Patent-Law Involving Chemistry." Harwood Huntington, 260 West Broadway, New York. This is a small pamphlet designed to show chemists some of the difficulties which attend the drafting of patents. 25 cents.

NEW PERIODICALS.

The Dyers' Bulletin, a semi-monthly journal devoted to practical dyeing, bleaching, printing, and finishing, and allied industries. The official organ of the Foremen Dyers' Mutual Improvement Association. Price, \$3 per year. 31 South Third Street, Philadelphia, Pa. This is a trade paper, and for all

such there is room, since the field is very large. Dyed samples are given, and the writing is addressed to the dyer.

The Hanna-Schoellkopf Company, of Buffalo, Philadelphia, and New York, issue a four-page trade paper entitled *Dye Topics*. It is a monthly publication devoted to the interests of the American dyehouses; incidentally, also, to the interests of American aniline colors, as sold by the Hanna-Schoellkopf Co. We are pleased to receive the paper, and wish it all success. There is room for all the trade papers, for the field is very vast. The newspaper is one of the most potent means of education, and should receive all encouragement.

A new periodical, the Revue Générale des Matières Colorantes, is commenced, and we have in our hands all of the issues of the first year. It would not be easy to commend this journal too strongly to color-chemists. It is not for the dyer, in the ordinary acceptance of the term, but is best adapted to the colorist who has had a training in chemistry. It is well printed, admirably furnished with dyed samples, and we place it in the first rank of all periodicals in the dyeing industry. Among the principal collaborators we find such names as Horace Koechlin, E. Noelting, Albert Scheurer, and Schaeffer, the acknowledged leaders in dyes and their application.

PART IV.--PATENTS.

No. 575,904—Black Azo Dye, Christopher R.s, assignor to John R. Geigy & Co., January 26, 1897.—A black powder soluble in water with bluish-black color, soluble in concentrated sulphuric acid with blue color, and which is transformed by complete reduction into a colorless solution.

No. 576,222—Rhodamin Dye, Carl Oscar Müller, assignor to Bindschedler, February 2, 1897.—Green, dyeing wool, silk and cotton a fine red, soluble in water, dissolving in alcohol with brick-red fluorescence, and dissolving in concentrated sulphuric acid and in hydrochloric acid with yellow coloration, which turns to red on adding water, the said dyestuff being precipitated from its aqueous solutions by an addition of hydrochloric acid.

No. 576,223—Rhodamin Dye, Carl Osear Müller, assignor to Bindschedler, February 2, 1897.—Green powder with metallic luster, dyes cotton, silk, and wool in a blue-red, dissolves in concentrated sulphuric acid with yellow coloration, and in water, alcohol, and ether with splendid red coloration.

No. 576,511.—Blue Trisazo Dye, Georg Steinike and Friedrich Schmidt, February 2, 1897.—Gray-black powder soluble in concentrated sulphuric acid with blue color, a violet precipitate on addition of water.

No. 578,093—Blue Dye, Albert Cobenzl, assignor to the Farbwerke, vormals Meister, Lucius & Brüning, March 2, 1897.—Brownish-black powder, soluble in water and alcohol with a pure blue color; in concentrated sulphuric acid with a green color, turning on addition of water first to a pure blue and then to a reddish blue; being soluble in ether, benzine, and ligroine, and separating from its aqueous solution on addition of a caustic alkali or ammonia blue flakes from its color base.

No. 578,432—Dark-Blue Azo Dye, Moritz Ulrich and Johann Nammann, assignors to the Farbenfabriken, vormals Bayer & Co., March 9, 1897.—A dark powder, soluble in concentrated sulphuric acid with a greenish-blue color, which turns, on the addition of ice-water to this sulphuric-acid solution first into bluish violet, and then darkish violet flakes are precipitated; on the addition of acetic acid (concentrated) to the hot watery solution of the dyestuff a violet solution results.

No. 578.578—Rhodamin Dye, Carl Oscar Müller, assignor to Basle Chemical Works Bindschedler, March 9, 1897.—Rhodamin dye, dyeing mordanted cotton with tannin and tartaricemetic. Soluble in water with red coloration, in alcohol with a yellowish fluorescence, and in concentrated sulphuric acid with a yellow coloration, turning to red on addition of water.

No. 578,580—Substantive Cotton Dye, Ferdinand Petersen, assignor to Ferd. Petersen & Co., March 9, 1897.—A black powder, dyes cotton; dissolves in concentrated sulphuric acid with a bluish-green coloration; gives, when hydrochloric acid is added to its aqueous solutions, a reddish-brown precipitate, and, when caustic soda is added to its aqueous solutions, a brown precipitate.

No. 579,773—Red-Blue Disazo Dye, Christian Rudolph, assignor to the firm of K. Oehler, March 30, 1897. A blackish-brown powder with a feeble metallic luster, soluble in water with a violet, and in concentrated sulphuric acid with a blue, color, and which dyes unmordanted cotton a red-blue.

No. 580.186—Blue Dye, Albert Herrmann, assignor to the Farbwerke, vormals Meister, Lucius & Brüning, April 6, 1897.

—A copper-colored powder of metallic luster, easily soluble in water, less easily in alcohol, its aqueous solution turning green with mineral acids, but not changing its color with ammonia or soda, turning violet on boiling with soda lye, and dyeing wool and silk a very even and fast blue in an acid bath.

No. 580,187—Fast Blue Dye, Albert Herrmann, assignor to the Farbwerke, vormals Meister, Lucius & Brüning, April 6, 1897. A copper-red powder of metallic luster, easily soluble in water, less easily soluble in alcohol, its aqueous solution turning green with mineral acids and not being changed by ammonia or soda, turning violet when boiled with soda-lye, and dyeing wool and silk a very even and fast blue in an acid-bath.

No. 580,188-Blue Acid Dye, Albert Herrmann, assignor to the Farbwerke, vormals Meister, Lucius & Brüning, April 6.

1897.—A copper-red powder of metallic luster, easily soluble in water, less easily soluble in alcohol, its aqueous solution turning green with mineral acids, but not changing its color with ammonia or soda, turning violet on boiling with sodalye, and dyeing wool and silk a very even and fast blue in an acid-bath.

No. 582,958—Trisazo Dye, Friedrich Schmidt and Otto Ernst, assignors to the Farbwerke, vormals Meister, Lucius & Brüning, May 18, 1897.—A black dyestuff, forms a grayish-brown powder soluble in water, with a blue-violet color. By adding ammonia the color turns a dull red-violet. It is soluble in concentrated sulphuric acid, with a blue color. By the addition of water the dyestuff is precipitated as gray-violet powder.

No. 582,959—Trisazo Dye, Friedrich Schmidt and Otto Ernst, assignors to the Farbwerke, vormals Meister, Lucius & Brüning, May 18, 1897.—A dyestuff, forms a violet-brown powder of metallic luster and is soluble in water with a bluish-violet color. The color of this solution changes on addition of ammonia into a dull red-violet. It is soluble in concentrated sulphuric acid with a blue color, while by addition of water a reddish-brown precipitate is formed.

No. 583,267—Blue-Green Dye, Jakob Schmidt, and Hugo Weil, assignors to Society of Chemical Industry in Basle, May 25, 1897.—A brown powder with a coppery luster, readily soluble in water, with a pure green-blue coloration, which does not change by the addition thereto of dilute acids.

No. 583,439—Black Azo Dye, Wilhelm Herzberg and Otto Hansmann, assignors to the Actien-Gesellschaft für Anilin Fabrikation, May 25, 1897.—A dark-brown powder of a metallic luster, soluble in hot water with a bluish-violet color, in which solution soda-lye produces a dark greenish-blue precipitate, acids precipitate the coloring acid as dark bluish-violet flakes, said dye being sparingly soluble in alcohol with a bluish-violet color, and in concentrated sulphuric acid with a blackish violet color.

No. 583,634—Black Trisazo Dye, Jakob Schmidt and Karl Jedlicka, assignors to the Society of Chemical Industry in Basle, June 1, 1897.—A dark powder of a light-bronze luster, capable of dyeing unmordanted cotton violet-black to greenblack, and which, when dissolved in concentrated sulphuric acid, imparts a blue coloration to the solution.

No. 583,635—Jakob Schmid and Karl Jedlicka, assignors to Society of Chemical Industry in Basle. June 1, 1897.—A powder of bronze luster, insoluble in alcohol, ether, and benzene, soluble in water with a dark blue to green coloration, and capable of dyeing unmordanted cotton in dark-blue to greenish-blue tints fast to light and to alkalies, and which, when dissolved in concentrated sulphuric acid, imparts a blue color to the solution, from which the dyestuff may be precipitated by an addition of water.

No. 584,119—Rhodamin Dye, Carl Oscar Müller, assignor to the Basle Chemical Works Bindschedler, June 8, 1897.—Fine green crystals, soluble in water and alcohol with a fine red tint, dissolve in concentrated sulphuric acid with a yellow tint, which turns to red on addition of water, and dyes cotton, silk, and wool in red tints.

No. 584,981—Blue Dye, Melchior Böniger, assignor to Chemische Fabrik vormals Sandoz, June 22, 1897.—A bronze powder, soluble easily in water with a pure blue shade, not altered by addition of soda, but turns reddish-violet on the addition of caustic soda, soluble in strong sulphuric acid, dyeing unmordanted cotton in a boiling-salt or Glauber-salt bath in bright-blue shades.

No. 585.104—Green Triazo Dye, Carl Jedlicka, assignor to the Society of Chemical Industry in Basle, June 22, 1897.—A powder of bronze luster, soluble in water with a green coloration, and capable of dyeing unmordanted cotton in darkgreen to yellowish tints, fast to light and especially to alkalies, and which, when dissolved in concentrated sulphuric acid, imparts a dirty blue-violet color to the solution.

No. 585,934—Blue Dye, Charles de la Harpe, assignor to L. Durand Huguenin & Co., July 6, 1897.—Dyes unmordanted wool and mordanted vegetable fibers, dissolves in concentrated sulphuric acid with a blue coloration, and is soluble in alkalies and alkaline carbonates with a blue-violet coloration.

No. 588,180—Bluish-Scarlet Dye, Ignaz Rosenberg, assignor to Kalle & Co., August 17, 1897.—Brownish-red bronzy powder, soluble in concentrated sulphuric acid with violet coloration, giving an alkaline reduction with zinc-dust, yellow solutions, which quickly turn brown under the influence of the air, dyeing on unmordanted cotton a bright bluish-scarlet.

No. 558.181-Reddish-Violet Dye, Ignaz Rosenberg, assignor

to Kalle & Co., August 17, 1897.—A black-bronze powder, easily soluble in hot water, insoluble in alcohol, soluble in concentrated sulphuric acid with blue coloration, giving, on alkaline reduction with zinc dust, a colorless solution, which turns brown under the influence of air, dyeing unmordanted cotton a bright reddish-violet.

No. 588,182—Mixed Substantive Dye, Ignaz Rosenberg, assignor to Kalle & Co., August 17, 1897.—A black-bronze powder easily soluble in hot water, insoluble in alcohol, soluble in concentrated sulphuric acid with blue coloration, giving, on alkaline reduction with zinc-dust, a slightly yellow solution, which turns brown when exposed to the air, dyeing unmordanted cotton a deep greenish-black.

No. 588,183—Brown Substantive Dye, Ignaz Rosenberg, assignor to Kalle & Co., August 17, 1898.—Dark-brown powder of greenish bronze, easily soluble in hot water, insoluble in alcohol, soluble in concentrated sulphuric acid with reddishviolet coloration, giving, on alkaline reduction with zinc-dust, colorless solutions quickly turning brown when exposed to the air, and dyeing a deep fast brown on unmordanted cotton.

No. 588,397—Gray Dye, Albert Cobenzl, assignor to the Farbwerke, vormals Meister, Lucius & Brüning, August 17, 1897.—A black powder easily soluble in water, difficultly soluble in alcohol, insoluble in ether, benzene, ligroin, etc., and soluble in concentrated sulphuric acid with a green color.

No. 589,766—Red Dye, Jules Ville, September 7, 1897.—Product being of a reddish color, soluble in alcohol and acetic acid, forming red solutions, and in concentrated hydrochloric and sulphuric acids, forming yellow solutions.

No. 590,088—Black Diazo Dye, Carl Billow, assignor to the Badische Anilin and Soda Fabrik, September 14, 1897.—Soluble in pure sulphuric acid, giving a reddish-blue to bluishgreen solution. On pouring into water, yields a blue flocculent precipitate, and on reduction in strong caustic alkaline solution, alpha-naphthylamin.

No. 591,309—Red Dye, Jules Ville, October 5, 1897.—Red coloring-matter resulting from heating a fatty hydrazin with rosolic acid, in hydrochloric and sulphuric acids forming yellowish solutions.

No. 591,616—Trisazo Dye, Melchior Böniger, assignor to Chemische Fabrik, vormals Sandoz, October 12, 1897.—Powders of slaty to purplish-brown color, dissolving in strong sulphuric acid with greenish-blue color, dyeing unmordanted cotton.

No. 592,608—Red Azin Dye. Bernhard Heymann, assignor to the Farbenfabriken of Elberfeld. October 26, 1897.—A brown powder, soluble in concentrated sulphuric acid with a green color, which turns into blue on the addition of a small quantity of ice-water to the sulphuric-acid solution, and finally from violet to red on the addition of a very large quantity of water, producing on mordanted cotton, as well as on unmordanted cotton, very clear red shades of great fastness to the action of alkali and light.

No. 593,347—Violet Disazo Dye, Fritz Bender, assignor to A. Leonhardt & Co., November 9, 1897.—Dark powders, dyeing cotton without mordant, yielding beautiful red to violet shades, and the solution in concentrated sulphuric acid having a greenish to bluish shade.

No. 593.790—Ulrich & Bammann, assignors to the Farben-fabriken of Elberfeld, November 16, 1897.—Powder with metallic, copper-like luster, soluble in cold water with violet-black color, soluble in alcohol with blue color, soluble in dilute ammonia liquor with blue color, soluble in concentrated sulphuric acid with greenish-blue color, which solution separates, on addition of a small quantity of ice-water, a greenish-blue, flaky precipitate, dyeing wool in acid-baths blue-black shades fast to the action of alkali and acid.

No. 594,105—Sulphur Dye, Henri Raymond Vidal, November 23, 1897.—Coloring-matter capable of directly dyeing cotton fibers a black or brown-black, soluble in sodium sulfid, appearing brown, being of a deep-brown color when unmixed with sodium sulfid.

No. 594,106—Mixed Sulphur Dye, Henri Vidal, November 23, 1897.—Coloring-matter directly soluble in acids, and dyeing animal fibers an intense black, and in part by a body soluble in alkalies, and dyeing vegetable fibers dull blue.

No. 594.107—Thiazin Dye, Henri Raymond Vidal, November 23, 1897.—Method of preparing coloring-matter which consists in heating with sulphur a mixture of paramidophenol and hydroquinone.

No. 594,123—Blue Tetrazo Dye, Christian Heinrich Rudolph and Julius Herbany, assignors to the firm of K. Oehler,

November 23, 1897.—Soluble in cold water, its aqueous solution not being altered by the addition of properly diluted caustic lye or soda, but yielding a red flocculent precipitate when the lye or soda is added in a concentrated condition, giving, when dissolved in concentrated sulphuric acid, a pure blue solution from which, by the addition of a little water, there are precipitated brown flakes which, by more water, are dissolved with a bluish-red color, dyeing cotton without mordants a bluish-red, which is distinguished by its pure, brilliant shade and fastness to acids.

No. 594,996—Black Azo Dye, Myrtil Kahn and Friedrich Runkel, assignors to the Farbenfabriken of Elberfeld, December 7, 1897.—A brownish-black powder, readily soluble in hot water with a violet-black color, in a twenty per cent. ammonia liquor with a bluish-black color, insoluble in alcohol, in a strong soda-lye and in concentrated hydrochloric acid, soluble in concentrated sulphuric acid with a black color, which turns into bluish-black on the addition of a small quantity of ice, while a violet-black precipitate is separated on the addition of a large quantity of ice to the sulphuric-acid solution, yielding on unmordanted cotton deep bluish shades, which are fast to acids, to alkalies, and to light.

No. 595,021—Black Azo Dye, Kahn & Runkel, assignors to the Farbenfabriken of Elberfeld, December 7, 1897.—Brownish-black powder, easily soluble in hot water, with a reddish brown-black color, and in caustic ammonia, with a violet-black color, insoluble in alcohol, in concentrated hydrochloric acid, and in strong soda-lye, soluble in concentrated sulphuric acid with a reddish violet-black color, which turns into violet-black on the addition of a larger quantity of ice to the sulphuric-acid solution, yielding, on unmordanted cotton, deep-black shades, which are fast to light, to acids, and to alkalies.

No. 595,349—Anthraufin Dye, Robert E. Schmidt, assignor to the Farbenfabriken of Elberfeld, December 14, 1897.—A dark-blue powder, readily dissolving in water with a pure blue color, from which solution it is precipitated by the addition of strong acids, soluble in diluted soda-lye with a pure greenish-blue color, in concentrated sulphuric acid with a yellow color, the color of the sulphuric-acid solution changing into greenish-blue on the addition of boric acid, the solution exhibiting in this state a characteristic absorption spectrum,

dyeing unmordanted wool in acid-baths intense pure and even shades, yielding on chromium mordants even greenishblue shades, which are fast to light and to fulling.

No. 595,350—Blue Dye from Chrysazin, Robert E. Schmidt, assignor to the Farbenfabriken, December 14, 1897.—A dark-violet powder with a copper-like luster, soluble in water with comparative ease, being precipitated from the watery solution by adding mineral acids, soluble in concentrated hydrochloric acid with a yellow color, from which solution it is precipitated by the addition of water, soluble in concentrated sulphuric acid with a yellow color, which is not changed on the addition of boric acid, soluble in alkalies with a pure greenish-blue color, insoluble in glacial acetic acid and in benzene, dyeing wool in acid-baths beautiful, even, and fast blue shades, and yielding, on chrome mordanted fibers, greenish-blue shades.

No. 596,333—Rhodol Dye, Carl Oskar Müller, assignor to the Basle Chemical Works, Bindschedler, December 28, 1897.—A dyestuff dissolving in water with a yellow-red color, and dyeing wool, silk, and tannin mordanted cotton of yellow-red tints, and which, in the form of its base, is soluble in concentrated sulphuric acid with a yellow color, and gives alkaline alcoholic solutions having a beautiful yellowish-green fluorescence.

No. 596,559—Brown Sulfuretted Dye, Arthur Weinberg, assignor to Leopold Cassella & Co., January 4, 1898.—Brown coloring-matter derived from dinitro-cresol, which is soluble in water in presence of sulfids or strong alkalies, insoluble in alcohol, neutral or acidulated water, the solutions of which dye unmordanted cotton directly brown.

No. 597,983—Black Substantive Cotton Dyestuff, Max H. Isler, assignor to the Badische Aniline and Soda Fabrik, January 25, 1898.—Black substantive coloring-matter for cotton, which can be derived from 1.4' orthodinitroanthraquinone and is soluble in cold water, giving a blue solution, and in concentrated sulphuric acid, giving a grayish-black color, slightly soluble in alcohol with green coloration, insoluble in dilute mineral acids, but soluble in alkalies and alkaline sulfids.

No. 599,425—Blue-Black Alizarin Dye, Robert E. Schmidt, assignor to the Farbenfabriken of Elberfeld Company, February 22, 1898.—A dark powder, difficultly soluble in cold water, readily soluble in hot water with a violet-red color,

soluble in dilute soda-lye with a dull violet-blue color, soluble in a dilute ammonia solution with a violet-red color, which turns into bluish-violet on the addition of an excess of ammonia liquor, easily soluble in hot alcohol with a violet-red color, which changes into violet-blue on adding an excess of ammonia liquor, and into greenish-blue on the addition of a small quantity of soda-lye, being decomposed under regeneration of purpurin when heated with a sulphuric acid of 60° Beaumé, yielding from bluish-gray to bluish-black, even shades on chromium mordanted wool, which shades are fast to light, to fulling, and against the action of acids.

No. 599,426—Green Dye derived from Anthraquinone, Robert E. Schmidt, assignor to the Farbenfabriken, February 22, 1898.—A dark-green powder, dissolving in water with a green color, which changes into yellow on acidulation and adding zinc-dust, the filtered solution reassuming the original green color on exposure to the air, and separating fine needles of quinizarin after heating with a small quantity of sulphuric acid, soluble in alcohol, acetone, chloroform, and benzene with a bluish-green color, soluble in concentrated sulphuric acid with a blue color, which changes into green on the addition of an excess of water, dyeing wool in acid-baths bluish-green shades, which withstand the action of light, of acids, and of alkalies in an excellent manner.

No. 599,427—Green Dye Derived from Quinizarin, Robert E. Schmidt, assignor to the Farbenfabriken of Elberfeld, February 22, 1898.—Green powder soluble in water with a green color, which changes into yellow on acidulating and adding zinc-dust, soluble in alcohol with a bluish-green color, in chloroform and acetone with a green color, in concentrated sulphuric acid with a bluish-green color, which changes into green on the addition of water, dyeing unmordanted wool in acid-baths green shades, which exhibit a great fastness to light, to the air, to acids, and alkalies.

No. 599,532 —Black Trisazo Dye, Christopher Ris, assignor to John R. Geigy & Co., February 22, 1898.—A black powder, soluble in water with black color, forming, with reducing agents, a colorless solution, soluble in concentrated sulphuric acid with a dark-blue color, scarcely soluble in alcohol, and dyeing unmordanted cotton, wool, and silk in deep black shades.

No. 601,033—Blue-Black Mixed Trisazo Dye, Melchior Böniger, assignor to the Chemische Fabrik, vormals Sandoz, March 22. 1898.—Coloring-matter nearly soluble in ethylic alcohol. Soluble in strong sulphuric acid with blue color, and forming, by reducing agents, benzidin, 1.4 naphthalenediamin, and easily soluble sulfo-acids of the naphthalene series.

No. 601,063—Benzidin-Orange, Christopher Ris, assignor to John R. Geigy & Co., March 22, 1898.—A brownish powder which is easily soluble in water with a deep orange color, scarcely soluble in alcohol, soluble in concentrated sulphuric acid with a red-violet color, precipitated by acids from its watery solution in brown flocks, and giving, when treated with reducing agents, benzidin or its homologues.

No. 601.363—Thiazin Dye, Henri Raymond Vidal, March 29, 1898.—Black powder, soluble in concentrated sulphuric acid, in sulfites, and soluble in hot alkalies generally.

No. 601.365—Black Dye, Henri Raymond Vidal, March 29, 1898.—Dark, porous mass, insoluble in acid, soluble in alkalies, giving a blue-black color, becoming darker on exposure to the air, and having the property of dyeing directly unmordanted cotton in shades of very deep black.

No. 601.\$59—Blue Azo Dye, Christian Rudolph, assignor to the firm of K. Oehler, April 5. 1898.—Black-violet powder, with a metallic luster, soluble in water with a blue-violet, and in sulphuric acid with a greenish-blue, color, muriatic acid changing the color of its watery solution into red-violet, and soda-lye into blue, zinc-dust decoloring its alkaline solutions.

No. 11.659—Black Dye and Process of Making Same, Raymond Vidal, April 5, 1898.—Dark mass, insoluble in water or acids, soluble in alkalies and alkaline sulfites, the alkaline solutions being readily oxidized on exposure to air, thereby becoming intensely black and capable of dyeing unmordanted cotton fiber in black shades.

No. 602,641—Brown Disazo Dye, Ernst König, April 19, 1898.—Blackish-brown powder, easily soluble in water with a brown color, which solution is not changed either by alkalies or acids, dissolved by concentrated sulphuric acid with an olive-green color, insoluble in ether and benzene or petroleum ether, but soluble in alcohol, and dyeing tanned and untanned cotton, as well as half-wool, brown in an acid-bath.

No. 602.640--Red-Violet Basic Disazo Dye, Ernst König, as-

signor to the Farbwerke, vormals Meister, Lucius & Brüning. April 19, 1898.—A brown powder, soluble in alcohol, ether, benzene, and petroleum ether, and dyeing tanned and untanned cotton, as well as half-wool, a reddish-violet in an acid-bath.

No. 602,639—Coppery-Brown Dye, Ernst König, assignor to the Farbwerke, April 19, 1898.—A black-green powder, easily soluble in water with a reddish yellowish-brown color, soluble in alcohol, insoluble in ether, benzene, and petroleum ether, and dyeing tanned and untanned cotton, as well as half-wool, a coppery brown.

No. 602,638—Basic Yellow Disazo Dye, Ernst König, assignor to the Farbwerke, vormals Meister, Lucius & Brüning, April 19, 1898.—An orange-yellow powder, soluble in water, alcohol, ether, and benzene, insoluble in petroleum ether, dyeing tanned and untanned cotton, as well as half-wool, yellow in an acid-bath.

No. 602,637—Basic Red Disazo Dye, Ernst König, assignor to the Farbwerke, vormals Meister, Lucius & Brüning, April 19, 1898.—A brown-red powder, soluble in water with a bluered color, soluble in alcohol, insoluble in benzene, ether, or petroleum ether, and dyeing tanned and untanned cotton, as well as half-wool, scarlet red in an acid-bath.

No. 602,544—Blue Azin Dye, Ott & Kroeber, assignors to the Farbenfabriken of Elberfeld Company, April 19, 1898.—Dark powder, soluble in water with a blue color, dissolving in concentrated sulphuric acid with a green color, which turns violet on the addition of a small quantity of ice, while a blue, flaky precipitate is separated on adding a larger quantity of ice, insoluble in a ten per cent. solution of sodium carbonate and in strong soda-lye, dyeing wool in acid-baths bright-blue, even shades, which are fast to acids and alkalies, and also pretty fast to the action of light.

No. 602,540—Violet Dye, Myrtil Kahn, assignor to the Farbenfabriken of Elberfeld Company, April 19, 1898.—Brownish-black powder, with a bronze-like luster, soluble in water with a brownish-violet color, in concentrated sulphuric acid with a blue color, which changes into bluish-violet on the addition of a small quantity of ice, while a blue, flaky precipitate is obtained on adding a larger quantity of ice to the sulphuric-acid solution, dyeing unmordanted cotton violet shades.

No. 603,093-Black Disazo Wool-Dye, Steinike & Schmidt,

assignors to the Farbwerke, vormals Meister, Lucius & Brüning, April 26, 1898.—A dark powder of bronze-like luster, easily soluble in water with a violet color, soluble with difficulty in alcohol with a reddish-blue color, and insoluble in benzene, the aqueous solution of which turns redder on the addition of ammonia, while in concentrated sulphuric acid the dyestuff is soluble with a pure blue color and is separated on addition of water as a reddish-violet precipitate, and dyes wool blue-black in an acid-bath.

No. 603,090—Brown Trisazo Dye, Krekeler & Martz, assignors to the Farbenfabriken of Elberfeld Company. April 26, 1898.—Dark-brown powder, soluble in water with a reddish-brown color, being dissolved by concentrated sulphuric acid with a blue color, dyeing cotton reddish-brown shades, which, when treated with solutions of chromium and copper salts, become more yellowish-brown and faster to washing and against the action of the light.

No. 603,016—Triphenylmethane-Blue Dye, Arnold Steiner, assignor to Chemische Fabrik, April 26, 1898.—Sulfonated blueviolet dyestuff, dissolving easily in concentrated sulphuric acid with a red-brown shade, in water and ethyl alcohol with a violet shade, and the aqueous solution of which, heated with caustic soda, turns first blue and then almost colorless, and may have the unchanged dyestuff precipitated from it by acids.

No. 603,013—Blue Azin Dye. Ott & Kroeber, assignors to the Farbenfabriken of Elberfeld Company. April 26, 1898.— A dark powder, soluble in water with a blue color, dissolving in concentrated sulphuric acid with a green color, insoluble in a teu per cent, solution of sodium carbonate and in strong soda-lye, dyeing wool in acid baths bright-blue shades, which are fast to acids and to alkalies, and also pretty fast against the action of light.

No. 603,009—Bluish-red Dye, Kahn & Runkel, assignors to the Farbenfabriken of Elberfeld Company, April 26, 1898.—A dark-brown powder, soluble in water, yielding a red solution, from which a black precipitate is obtained on the addition of dilute hydrochloric or sulphuric acid, soluble in concentrated sulphuric acid with a bluish-black color, which turns violet on the addition of a small quantity of ice, while a black, flaky precipitate is obtained on the addition of a larger quantity of ice to the sulphuric-acid solution, yielding, on unmor-

danted cotton, intense bluish-red shades, which, when treated with a solution of diazotized paranitranilin and sodium acetate, change into intense brown shades fast against washing.

No. 603,008—Violet Azo Dye, Myrtil Kahn, assignor to the Farbenfabriken of Elberfeld Company, April 26, 1898.—Black powder of a bronze-like luster, soluble in water with a violet color, and in concentrated sulphuric acid yielding a blue solution, from which a blue, flaky precipitate is obtained on the addition of a sufficient quantity of ice, dyeing unmordanted cotton violet shades, which, when treated with solutions of diazotized paranitranilin and sodium acetate, change into black shades fast to acids and to washing.

No. 602,857—Black Trisazo Dye, Krekeler & Israel, assignors to the Farbenfabriken of Elberfeld Company, April 26, 1898.—A dark powder, dissolving in water with a violet-gray color, soluble in concentrated sulphuric acid with a blue color, yielding on unmordanted cotton violet-black shades, which, when treated with solutions of chromium and copper salts, change into greenish-black fast to washing.

No. 602,858—Brown Trisazo Dye, Krekeler & Martz, assignors to the Farbenfabriken of Elberfeld Company, April 26, 1898.—Dark-brown powder, soluble in water with a yellowish-brown color, in concentrated sulphuric acid with a blue color, dyeing unmordanted cotton yellowish-brown shades, which, when treated with solutions of chromium and copper salts, become more reddish and faster to washing and against the action of light, yielding on wool in acid-baths yellowish-brown shades, which, on a treatment with chromium salts, become fast against milling.

No. 602,855—Blue-black Trisazo Dye, Krekeler & Blank, assignors to the Farbenfabriken of Elberfeld Company, April 26, 1898.—Dark powder with bronze-like luster, dissolving in water with a blue color, in concentrated sulphuric acid with a greenish-blue color, yielding on reduction with stannous chloride in a hydrochloric-acid solution a mixture of reduction products from which tolidin and a diamidonaphthosulfo acid can easily be isolated.

No. 602,856—Black Trisazo Dye, Krekeler & Israel, assignors to the Farbenfabriken of Elberfeld Company, April 26, 1898.—A dark powder dissolving in water, yielding blue solutions, soluble in concentrated sulphuric acid with a blue color,

dyeing unmordanted cotton blue shades, which, when treated with solutions of chromium and copper salts, change into black fast to washing.

No. 603,659—Dark-green Alizarin Derivative, Robert E. Schmidt, assignor to the Farbenfabriken of Elberfeld Company, May 10, 1898.—Dark-green powder, dissolving in water with a green color, yielding, on chrome-mordanted cotton, green shades which are distinguished by their brightness and fastness against the action of light.

No. 603.645—Green Trisazo Dye, Krekeler & Martz, assignors to the Farbenfabriken of Elberfeld Company, May 10. 1898.—A black powder, soluble in water with a green color, in concentrated sulphuric acid with a dark-blue color, from which solution a greenish-black precipitate is obtained on the addition of a sufficient quantity of ice, dyeing unmordanted cotton green shades.

No. 603.646—Gray Trisazo Dye, Krekeler, Martz & Israel, assignors to the Farbenfabriken of Elberfeld Company. May 10, 1598.—A dark-brown powder, soluble in water with a greenish-gray color, in concentrated sulphuric acid with a blue-black color, from which a black, flaky precipitate is obtained on the addition of a sufficient quantity of ice, dyeing unmordanted cotton greenish-gray shades.

No. 603,647—Green Trisazo Dye, Krekeler, Martz & Israel. assignors to the Farbenfabriken of Elberfeld Company. May 10, 1898.—A black powder, soluble in water with a green color, which, on the addition of alkalies, turns into greenish-black, in concentrated sulphuric acid with a blue-black color, from which solution a greenish-black, flaky precipitate is obtained on the addition of a sufficient quantity of ice, dyeing unmordanted cotton green shades.

No. 603,648—Brown Trisazo Dye, Krekeler, Martz & Israel, assignors to the Farbenfabriken of Elberfeld Company, May 10. 1598.—A dark-brown, nearly black powder, soluble in water with a violet-brown color, in concentrated sulphuric acid with a bluish-black color, from which solution a violet-black precipitate is obtained on the addition of a sufficient quantity of ice, dyeing unmordanted cotton violet-brown shades, which, when treated with solutions of chromium and copper salts, become a dark brown with a yellowish hue, and faster to washing and against the action of light, yielding, on wool in acid-baths, dark-brown shades.

No. 603,755—Brown Sulphur Dye, Robert Demuth, assignor to the Farbenfabriken of Elberfeld Company, May 10, 1898.— A brownish-black powder insoluble in water, sodium carbonate, ammonia, and alcohol, soluble with difficulty in dilute soda-lye at ordinary temperature, more easily soluble in hot soda-lye with a greenish-brown color, insoluble, or soluble with difficulty, in concentrated sulphuric acid, easily soluble with a greenish-brown color in alkaline sulfids, thus affording solutions which dye cotton at ordinary temperatures in fast brown shades.

No. 605,119—Diphenylnaphthylmethane Dye, Osear Nastvogel, assignor to the Farbenfabriken of Elberfeld Company, June 7, 1898.—A brown powder, easily soluble in water with a greenish-blue color, in concentrated hydrochloric acid with a yellow color, in concentrated sulphuric acid with a yellowish color, which changes into green on the addition of a sufficient quantity of ice, being partly dissolved with a reddishviolet color under evolution of dimethylamin when heated with a strong soda-lye, dyeing wool in acid-baths clear, greenish-blue, and even shades fast to alkalies.

No. 605,103—Black Disazo Dye and Process of Making Same, Kahn & Heidenreich, assignors to the Farbenfabriken of Elberfeld Company, June 7, 1898.—A brownish-black powder, soluble in water with a bluish-black color, in concentrated sulphurie acid with a greenish-black color, which changes into violet-black on adding a small quantity of ice, while a violet-black precipitate is obtained on the addition of a larger quantity of ice to the sulphurie-acid solution, dyeing wool in acid-baths black shades fast to milling and against the action of light.

No. 605,568—Anthraquinone Dye, Robert E. Schmidt, assignor to the Farbenfabriken of Elberfeld Company, June 14, 1898.—A blackish powder, which, when rubbed, assumes a metallic luster, insoluble in alcohol, ether, benzene, nitrobenzene, and in glacial acetic acid, soluble in anilin and pyridin with a pure greenish-blue color, in dilute alkalies with a blue color, yielding, with concentrated sulfiburic acid a yellow solution, which assumes a blue color on the addition of boric acid, and shows in this state a characteristic absorption spectrum, dyeing unmordanted wool in acid-baths beautiful blue shades fast to light, yielding on chrome-mordanted wool greenish-blue shades very fast to light.

No. 605,921—Blue Anthraquinone Dye, Schmidt & Tust, assignors to the Farbenfabriken of Elberfeld Company, June 21, 1898.—A blackish powder, which, when rubbed, assumes a metallic luster, soluble in anilin, pyridin, and in dilute alkalies with blue color, yielding, with concentrated sulphuric acid, a yellow solution, which assumes a violet-red color on the addition of boric acid, dyeing unmordanted wool in acidbaths beautiful blue shades very fast to light, yielding, on chrome-mordanted wool, greenish-blue shades very fast to light and milling.

No. 606,436—Blue Dye, Fritz Bender, assignor to Farbwerk Mühlheim, vormals A. Leonhardt & Co., June 28, 1898.—Dark powder with metallic luster, soluble in water, giving a pure blue solution, insoluble in spirit, soluble in concentrated sulphuric acid with a pure blue color, and dyeing unmordanted cotton a pure blue shade.

No. 606,181—Blue Tetrazo Dye and Process of Making Same, Johann Bammann, assignor to the Farbenfabriken of Elberfeld Company. June 28, 1898.—A dark powder with a bronzy luster, easily soluble in water with a reddish-blue color, being dissolved by concentrated sulphuric acid with a greenish-blue color, which turns into bluish violet on adding a small quantity of ice, while a bluish-violet, flaky precipitate is obtained on the addition of a larger quantity of ice to the sulphuricacid solution, dyeing unmordanted cotton violet-blue shades fast to acids and alkalies.

No. 506.193—Yellow-brown Cotton Dye, Robert Demuth, assignor to the Farbenfabriken of Elberfeld Company, June 28, 1898.—A dark-brown mass or powder, readily soluble in water with a reddish-brown color, unalterable by the addition of sodium sulphide, being separated from its watery solution by the addition of dilute hydrochloric or sulphuric acid as a brown precipitate, while sulphuretted hydrogen is set free, yielding on unmordanted cotton, in cold or hot baths, with or without the addition of alkaline sulphid, bright yellowish-brown shades fast against the action of alkalies and light.

No. 606.212—Blue Dye and Process of Making Same, Bernhard Heymann, assignor to the Farbenfabriken of Elberfeld Company, June 28, 1898.—A dark powder, soluble in hot water with a blue color, in concentrated sulphuric acid with a brown color, which changes into yellowish-green on the addition of

a small quantity of ice, soluble in concentrated hydrochloric acid with a yellowish-green color, from which solution a violet, flaky precipitate is obtained on the addition of water, dyeing chrome-mordanted wool bright-blue shades fast against light and milling.

No. 606,264—Dark-blue Tetrazo Dye and Process of Making Same, Johann Bammann, assignor to the Farbenfabriken of Elberfeld Company, June 28, 1898.—A dark powder with a metallic luster, easily soluble in water with a blackish-blue color, in concentrated sulphuric acid with a pure blue color, which changes into violet on the addition of a small quantity of ice to the sulphuric-acid solution, being rediazotizable in solution or when on fiber, yielding on unmordanted cotton blackish-blue shades.

No. 606,295—Blue Azin Dye and Process of Making Same, Ott & Kroeber, assignors to the Farbenfabriken of Elberfeld Company, June 28, 1898.—A darkish-brown powder, soluble in water, with a splendid blue color, in concentrated sulphuric acid with a green color, which turns into blue on mixing the sulphuric acid with a large quantity of water, dyeing wool brilliant blue shades fast to the action of alkalies, of acids, and likewise fast against milling.

No. 606,439—Green Disazo Dye, Fritz Bender, assignor to Farbwerk, Mühlheim, vormals A. Leonhardt & Co., June 28, 1898.—A dark powder, dissolving in water with a green color, being scarcely soluble in spirit, giving, with concentrated sulphuric acid, a violet solution, which, when diluted with water, yields a violet precipitate, said coloring-matter dyeing unmordanted cotton in green shades.

No. 606,438—Blue-black Dye, Fritz Bender, assignor to Farbwerk Mühlheim, June 28, 1898.—Dyestuff soluble in water with a blue color, but scarcely soluble in spirit, and the solution of which in concentrated sulphuric acid has a bluishgreen tint, and when diluted with water yields a blue precipitate, the said dyestuff dyeing wool in bluish-black shades.

No. 607,408—Blue Dye for Wool, Emil Elsaesser, assignor to Dahl & Co., July 12, 1898.—A dark-bronze, shining powder, which is readily soluble and takes on a blue color in water, but is with difficulty soluble in alcohol, and, with the addition of hydrochloric acid to the aqueous solution, is rendered somewhat greener, while the addition of soda-lye makes it somewhat redder.

No. 608,024—Brown Azo Dye, Melchior Böniger, assignor to Chemische Fabrik, vormals Sandoz, July 26, 1898.—A brownblack powder, sparingly soluble in ethylic alcohol, easily soluble in hot water with brown color, turning in intense redbrown on addition of caustic soda, soluble in concentrated sulphuric acid with blue-violet color, giving a bluish-brown precipitate on addition of water, and dyeing unmordanted cotton in yellow-brown to blue-brown tints.

No. 608,238—Green Alizarin Dye, Karl Thun, assignor to the Farbenfabriken of Elberfeld Company, August 2, 1898.—A dark-green powder, dissolving in water with a green color, which changes into yellow on acidulating and adding zincdust, the filtered solution separating leuco-alizarin bordeaux after heating with a small quantity of sulphuric acid, dyeing unmordanted and chrome-mordanted wool green shades fast to light, yielding on chrome-mordanted cotton green shades, which are distinguished by their brightness and fastness against the action of light.

No. 608,355—Brown-black Sulphur Dye, Henri Raymond Vidal, August 2, 1898.—A dark mass, soluble in alkalies, insoluble in acid, and having the property of directly dyeing unmordanted cotton a brown black.

No. 609,598—Red Dye and Process for Making Same, Henri Raymond Vidal, August 23, 1898.—A vivid red-blue mass, soluble in alkalies, capable of dyeing directly animal and mordanted cotton fibers.

No. 609,131—Mordant, Gottlob Wendler, August 16, 1898.— Mordant, consisting of or containing 60 parts, by weight, of commercial sulphuric acid, 100 parts of calcined alkaline sulphate, such as sodium sulphate, and 15 parts of boracic acid.

No. 608,999—Blue-Black Azo Dye, Bammann & Ulrich, August 16, 1898.—A brownish-black powder of metallic luster, soluble in water and alcohol with reddish-violet color, in ammonia with bright reddish violet, slightly redder than the aqueous solution, little soluble in sodium carbonate at ordinary temperature, more on heating, with a dull reddish-violet color, from which solution brownish-black flakes separate on cooling, searcely soluble in soda-lye at ordinary temperature, more on heating, with a dull violet color, almost insoluble in dilute hydrochloric and sulphuric acid, even on heating, soluble in concentrated sulphuric acid with a greenish-blue color,

assuming, on addition of ice-water, first, a pure blue, then a reddish-blue color, and finally separating a fine blue precipitate, dyeing unmordanted cotton in an alkaline bath violet-black shades, which, after treatment with nitrous acid and subsequent combination with an alkaline solution of betanaphthol, change into black.

No. 609,327—Blue-black Dye and Process of Making Same, René Bogn, assignor to the Badische Anilin and Soda Fabrik, August 16, 1898.—A new coloring-matter, which can be derived from dinitronaphthalene, and which is soluble in water, gives a blue color in ammoniacal water, and which yields a fluorescent bluish or bluish-green color in concentrated sulphuric acid.

No. 609,352—Blue Dye, Paul Julius, assignor to the Badische Anilin and Soda Fabrik, August 16, 1898.—Blue coloring-matter, readily soluble in water, giving pure blue solutions that are hardly changed in color by the addition of an alkali, and giving a dirty reddish-brown color approaching black in concentrated hydrochloric acid, and a reddish-brown color in concentrated sulphuric acid.

No. 609,998—Sulfonated Monobenzyl-Phthalein Dye, Schmid & Rey, assignors to the Badische Anilin and Soda Fabrik, August 30, 1898.—Dye-stuff which is the alkaline sulfonate of monobenzylated-phthalein dye, dyeing textile fibers in fiery red tints, and being a red powder, which is soluble in water with a red coloration and a yellowish fluorescence, with difficulty soluble in alcohol, and insoluble in ether and benzene.

No. 609,997—Red-violet Dye, Schmid & Rey, assignors to the Badische Anilin and Soda Fabrik, August 30, 1898.—A dye-stuff which dyes wool and silk in acid-bath in fine red-violet tints, dissolves in water with a blue-red color and a yellowish fluorescence, is sparingly soluble in alcohol, insoluble in ether and benzene, dissolves in concentrated sulphuric acid with yellow coloration, and may be precipitated from its solution in concentrated sulphuric acid, by an addition of water, in the form of free sulfo-acid.

No. 610,367—Basic Disazo Dye, Alfred Philips, assignor to the Farbwerke, vormals Meister, Lucius & Brüning, September 6, 1898.—A red powder, soluble in water and alcohol with a red color, soluble with difficulty in benzene, insoluble in ether and petroleum ether, and dyeing wool and cotton cherryred in an acid bath. No. 610.349—Violet Azo Dye, Otto Ernst, assignor to the Farbwerke, vormals Meister, Lucius & Brüning, September 6, 1898.—A green-black powder of metallic luster, soluble with difficulty in cold, more easily in hot water, with a red-violet color, soluble in concentrated sulphuric acid with a graygreen color, and giving, by the addition of water, a violet precipitate, dyeing wool violet in an acid-bath, and producing, by treatment with chromates or chromic acid, an even, saturated brown color of beautiful shade, fast to milling and light.

No. 610,345—Red Acid Dye, Bernhard Deicke, assignor to the Farbwerke, vormals Meister, Lucius & Brüning, September 6, 1895.—A dark powder, soluble in water and alcohol with a red color, almost insoluble in ether, benzene, and ligroin.

No. 610,541—Black Dye, George Kalischer, assignor to Cassella & Co.—A black mass, easily soluble in water with a blueblack color, being precipitated from its solutions by acids, and dyeing unmordanted cotton directly in an alkaline bath, without the help of oxidizing agents, a deep blue-black.

PART V.—NEW DYES.

SYNTHETIC INDIGO.

The Société Industrielle de Mulhouse has granted two medals of honor in connection with the recent production of artificial indigo. One of these medals was given to the Badische Anilin & Soda Fabrik, with the words, "The Industrial Society rarely has had occasion to give a prize so well merited, and rarely have the conditions been fulfilled in a manner so complete; upon this point there has been one voice in the Committee of Chemistry."

The other medal of honor has been given to M. Von Baeyer, and it is the prize known as the medal of honor for the synthesis of one of the natural coloring principles which has an industrial application. The words in connection with this, "The Society thanks you at one and the same time for your work in pure and applied science." It is with great difficulty that one unfamiliar with the enormous amount of work required in the synthesis of indigo appreciates the beauty of the successful production of indigo. It is true that there have been hitherto sold on the market compounds which under certain conditions could be used for the making of an indigo. Two of these, which were on the market for some time, demanded the use of subsequent chemicals to fully develop the color. To-day the Badische Anilin & Soda Fabrik sells an indigo which on analysis gives 98 and 99 per cent. of pure indigotin. Moreover, it is a constant composition, which is a fact of vast importance to those using indigo.

FARBENFABRIKEN OF ELBERFELD.

Acid Violet 3 B extra. Acid Violet 4 B extra. Acid Violet R extra. Acid Violet 2 R extra.

Acid Violet 3 R extra.

Acid Violet RX.

Acid Violet 3 BX.

Acid Violet 4 BX.

Alizarine Green CG extra.

Alizarine Green CK.

Alizarine Viridine Paste.

Alizarine Green CE Paste.

Alizarine Brown RB.

Benzo Chrome Black Blue.

Benzo Chrome Black Blue B.

Benzo Green G.

Brilliant Alizarine Blue D Paste.

Diamond Flavine G.

Diazo Black 3 B.

Diazo Black BHN.

Double Ponceau 2 R, 3 R and 4 R.

Phenol Black SS.

Pluto Black G.

Pluto Black R.

Sulphon Acid Blue B.

Sulphon Acid Blue R.

KALLE & CO.

Biebrich Alizarine Black 4 B N patented.

Biebrich Acid Blue G patented.

Heliotrope 2 B.

Naphtamine Blue 2 B patented.

Naphtamine Blue 2 R patented.

Naphtamine Blue 3 R patented.

Naphtamine Blue 5 B patented.

Naphtamine Indigo Blue 2 B patented.

Naphtamine Indigo Blue 2 R., patented.

Naphtamine Indigo Blue G patented.

Salicine Yellow.

Tolan Red B patented.

JOHN J. KELLER & Co.

Acid Grenadine.
Acid Grenadine B.

Diphenyl Black R.
Diphenyl Blue E F S.
Diphenyl Blue Black.
Diphenyl Brown.
Diphenyl Brown R.
Diphenyl Brown R R.
Diphenyl Fast Black.
Diphenyl Fast Yellow G.
Diphenyl Indigo Blue.
Diphenyl Red.
Diphenyl Violet R.
Diphenyl Yellow.
Eriocyanine.
Erioglaucine.

VICTOR KOECHL & Co.

Alizarine Black R G (Cotton).
Alizarine Black D R.
Alizarine Blue D G.
Alizarine Blue D R (Cotton).
Azophor Black S (Printing).
Azophor Orange (Printing).
Blue Rongeant, P P W (Printing).
Cœrulein B W paste.
Dianil Black G.
Dianil Black R.
Dianil Blue B G.
Dianil Blue R.
Fast Silk Gray O (Silk).
Also the series of Janus Colors.

'NEW YORK & BOSTON DYEWOOD COMPANY.

Brilliant Bordeaux S mixture.
Columbia Brown R.
Columbia Black F B.
Chromanil Black R F.
Chromanil Brown G G and R.
Sambesi Black B R.
Sambesi Black D and 3 B.
Wool Black 4 B F.

Wool Black B.
Wool Black G R.
Wool Jet Black 2 B and 3 B.

WILLIAM J. MATHESON & Co., LTD.

Alizarine Black R patented.

Alizarine Black S patented.

Alizarine Black T patented.

Alizarine Lanacyl Navy Blue B patented.

Anthracene Acid Black S T.

Anthracene Acid Brown N.

Anthracene Acid Brown S W, patented.

Bismarck Brown-Yellow Shade.

Diamine Azo Black B.

Diamine Azo Black B patented.

Diamine Azo Blue, No. 51, patented.

Diamine Azo Blue, No. 54, patented.

Diamine Azo Blue, 72,122, patented.

Diamine Azo Blue R R.

Diamine Black B.

Diamine Blue, No. 50, patented.

Diamine Blue, No. 52, patented.

Diamine Blue, No. 53, patented.

Diamine Blue, No. 55, patented. Diamine Blue S R X, patented.

Diamine Blue Black R.

Diamine Brown 2 O patented.

Diamine Brown, No. 30a, patented.

Diamine Brown, No. 31, patented.

Diamine Brown, No. 32, patented.

Diamine Brown, No. 33, patented.

Diamine Brown, No. 34, patented.

Diamine Brown, No. 35, patented. Diamine Brown, No. 36, patented.

Diamine Brown, No. 37, patented.

Diamine Catechine G patented.

Diamine Catechine B.

Diamine Nitrazol Brown B patented.

Diamine Nitrazol Brown B D patented.

Diamine Nitrazol Brown G patented.

Diamine Nitrazol Brown R D patented.

Diamine Nitrazol Black B.

Diamine Red 10 B.

Diamineral Blue R patented.

Diaminogen B R.

Excelsior Lake Scarlet J N.

Fast Acid Green B.

Formyl Violet 8 B patented.

Gloria Black B patented.

Immedial Black V extra patented.

Lake Searlet F R.

Lake Searlet F R R.

Lake Scarlet F R R R.

Lanafuchsine S B.

Lanafuchsine S G.

Milling Yellow I I.

Naphtindone T patented.

Oxy-Diamine Black A patented.

Oxy-Diamine Black A M patented.

Oxy-Diamine Black A T patented.

Oxy-Diamine Black B G patented.

Oxy-Diamine Black R R patented.

Oxy-Diamine Blue 3 R patented.

Oxy-Diamine Orange G patented.

Oxy-Diamine Orange R patented.

Oxy-Diamine Yellow G G patented.

Scarlet E C.

Wool Red B.

WILLIAM PICKHARDT & KUTTROFF.

Alizarine Blue N A extra.

Alizarine Blue N G G.

Alizarine Yellow L W.

Cotton Black B.

Neptune Green.

Nile Blue R.

Palatine Black.

Palatine Scarlet.

Sorbin Red.

Union Black B B.

Wool Blue.

READ HOLLIDAY & SONS, LTD.

Acid Black, 5,534 and 5,535.

Cross Dve Black.

Cross Dye Drab.

Naphthol Black 4 B.

Titan Blue 3 B.

Titan Blue R.

Titan Brown R.

Titan Como S. Titan Navy R.

Titan Red.

Titan Red 6 B.

Titan Sky Blue.

Titan Yellow R.

Titan Yellow Y.

X. L. Blue.

X. L. Red.

Vat Red Paste.

Vat Red Powder.

SYKES & STREET.

Brown S. D. M.

Brown S. D. P.

Induline 3 B.

Rouge de Lyon A.

Soudan Black, 1, 2, and 3.

Soudan Black B.

Sondan Black R.

Soudan Black 2 R.

Thio Catechine, Nos. 2, 3, and 4.

Violet A P F.

Violet Blue A P.

PART VI.—NEW PROCESSES.

COP-DYEING.

The real key to the difficulty is in matching lots dyed at different times. To a layman nothing appears easier than to obtain the same color by taking precisely the same amounts of chemicals and the same weight of goods, and in this way try to obtain uniform dyeings, day after day, and week after week. It is a fact, however, that no one who has had any experience in colors will gainsay, that it is impossible, even where the most conscientious care is used. to get identically the same results day in and day out. All of this seems absurd to the theorist, but it is a sad fact that the dyer cannot get uniform shades even when using, to the very best of his ability, identically the same amounts. Therefore, as the order always comes from the commission house, "Dye THIS red," and not "Dye A red," it is not possible to make a commercial success on a commission basis of a process which does not enable the dyer to return the goods to the commission house dyed precisely and identically THIS red. Per contra, if a mill uses up its own dyeings, the dyeing of the cops can be used to effect a very material economy.

Another cause for a great deal of trouble in carrying out to a practical success the idea of cop-dyeing is that, if the cops are handled in the least roughly or squeezed so that the shuttling is in the least delayed, there is a loss, because in this country, where several looms are kept in operation by one hand, this operation of shuttling must be easily done, or the whole bobbin is thrown in the waste heap. We have not the careful labor in this country, and it is made more careless because of the migration of mill help from one manufacturing center to another. Abroad this migration is not so easy, and consequently the mill hands stay longer in one place and get used to a certain kind of work.

To enumerate cop-dyeing machines: The first, that of M. Charles Weber; second is a cop-dyeing machine used in Providence; and third is mentioned in the Consular Reports for September, 1892, page 152; Mr. C. R. English, of Boston, is agent, we believe, for more than one cop-dyeing machine, and should be addressed.

Perhaps the newest machine in cop-dyeing is the Hallewell machine, for which R. B. Maclea, \$3 Worth Street, New York City, is the agent.

A late number of the Textile Manufacturer has an article on an improved cop-dyeing machine made by Mr. Beaumont, of Stockport.

DYEING IN SLUBBING AND ROVING.

An English patent has been taken for bleaching and dyeing in the various stages of manufacture. The idea is to use perforated bobbins placed on the frames, and these specially constructed bobbins are so arranged that they screw on to large pipes, through which come bleach and dye liquors. The idea is old.

SLIVER DYEING MACHINES.

A very handsome announcement appears in a recent issue of the Textile Manufacturer of a continuous sliver dyeing machine, manufactured by the firm of Mather & Platt. The idea is to pass the slivers six at a time, between perforated bands, which prevent disturbance of the fibers. These bands come together over the slivers, and then bands, slivers, and all pass through the different dyeing solutions, either hot or cold. The whole machine shows an enormous amount of labor, thought, and enterprise. It is a matter of regret to say that we do not think there is anything in it, so far as commission work is concerned.

PATENT DYEING PROCESSES.

Sprague's patent dyeing apparatus for dyeing raw stock gives excellent results. The first experiments have completely discouraged some, but pertinacity will make the process do good work.

The Bentz-Edmeston continuous kier has been favorably

reported upon by some firms using it. The process is based on the employment of caustic soda in a machine.

The electrical bleaching process of Gebaur-Knofler has been used in Berlin. The theory is the electrolysis of a solution of common salt.

CALICO PRINTING-DOUBLE WIDTH.

The Scotch printers have been early in the field with a method for double printing, and the production is very economical, being double what the ordinary machine will turn out. The process is known as the "split" trade. A machine printing "splits" is one that turns out the double width of cloth, printing practically two pieces at one time. In order to do this, the machine is made only twenty inches wider than the ordinary one. The owners of the machines that print but one width are practically obliged to take 50 per cent. of their previous prices in order to compete with the mills owning "split" machines.

COUPLING PROCESS.

Every cotton dyer is familiar with the now well-known process of dyeing and then, in subsequent baths, diazotizing with sodium nitrite and developing with such substances as naphthol, phenylene diamine, para-nitraniline, etc.

There has been placed on the market "Nitrazol C," a ready prepared form of diazotized para-nitraniline, which only requires dissolving in water to be ready for use. To distinguish this process from other methods of dyeing cotton, it may be very conveniently called the "Coupling Process."

In working with para-nitraniline: Take 2 pounds paranitraniline C. and dissolve in 1½ gallons of boiling water and ½ gallon of hydrochloric acid. Stir until a complete solution is obtained, then add 3½ gallons of cold water, and allow the mass to stand until thoroughly cold; add 1½ pound of nitrite of soda dissolved in 2/3 gallon of cold water. In from fifteen to twenty minutes the diazotization of the para-nitraniline will be complete, when sufficient cold water is added to make up to 20 gallons. This is kept as a stock solution, and if kept in wood or earthen vessels, not exposed to heat or light, will keep for two or three weeks. To make the coupling bath, take, for 100 pounds of cotton, 10 to 20 gallons; mix with enough water to work the cotton in handily, and add for every 10 gallons of the stock liquor used 2 pounds of acetate of soda. The first-named quantity is used for producing dark shades.

With Nitrazol C.: For 100 pounds of cotton, take from 4 to 8 pounds Nitrazol C. and dissolve in cold water: add, for each pound of Nitrazol used, ½ pound of refined alkali and 1½ ounce acetate of soda, both previously dissolved in water. When light shades are being produced, 4 to 5 pounds of the Nitrazol will be found sufficient; dark shades will take from 6 to 8 pounds.

Not all of the Diamine Colors are suitable for this coupling process, the following being those best adapted: Diamine Jet Black OO, patented; Diamine Jet Black CR, patented; Diamine Jet Black RB, patented: Oxy-Diamine Black, SOOO, patented; Diamine Brown V. patented; Cotton Brown N, Cotton Brown A, Diamine Blue Black E. patented; Diamine Black BO, patented; Diamine Gray G, patented; Diamine Gray G, patented; Primuline, Diamine Orange D, Diamine Bronze G. patented; Primuline, Diamine Orange D, Diamine Fast Yellow A; Diamine Brown M, patented: Diamine Black RO, patented; Diamine Nitrazol Brown Y, patented; Diamine Nitrazol Brown BD, patented; Diamine Nitrazol Brown GD, patented; Diaminogene B, patented.

DEVELOPERS.

PHENYLENE DIAMINE.

This is a product of almost chemical purity, manufactured especially for the production of fast dyeing with Diamine Colors, and is used for the development of blacks with Diamine Blacks RO, BO, BH, Diamine Jet Black SS, Diamine Blue B'ack E. Diaminogene Extra, D'aminog ne B, and Diamine Black NB, and for the development of browns with Diamine Cutch, Diamine Browns M and V. One-half per cent, of phenylene diamine is sufficient for the development of full shades.

NAPHTHYLAMINE ETHER N, PATENTED.

This product is used for the development of fast dyeings with the Diamine Colors; the shades produced tend toward blue tones, and it is especially valuable for the production of bright blues. It is used chiefly for the production of blues from Diamine Blacks RO, BO, BH, Diamine Blue Black E, Diaminogene Extra, Diaminogene B, Diamine Azo Black B and R. The blues thus produced are unsurpassed in fastness and brilliancy. About 1 per cent. is sufficient for average shades.

FAST BLUE DEVELOPER AD, PATENTED.

This developer gives shades with the Diamine Colors, similar to those produced with naphthylamine ether N, but of a somewhat redder tone. It has the additional value of producing fast shades with the Diamine Blues. The same quantity is necessary as with naphthylamine ether N.

BLUE DEVELOPER AN, PATENTED.

Very similar in results to naphthylamine ether N.

NITRAZOL C, PATENTED.

This product is diazotized paranitraniline in a dry, powdered form. It is stable at a temperature below 85° F., and is well adapted for use in dye-houses where facilities for diazotizing cannot be obtained. It is liable to decomposition at a temperature of about 85° F., and should be kept in a cool, dry place. For use, 5 per cent. Nitrazol C are dissolved in cold water, and 1/10 per cent. of sodium acetate and ¼ per cent. of sodia ash, of the weight of the Nitrazol, are added. The goods, dyed with the proper Diamine Colors, are worked in this cold solution. This method is used chiefly in producing browns when coupled with Diamine Jet Blacks OO, RB, CR, Oxy-Diamine Blacks SOOO, Diamine Brown V, and Cotton Browns N and A. Also with Diamine Blue Black E, Diamine Black BO, Diaminogene, Primuline, Diamine Orange D, and Diamine Fast Yellow A.

AZOTOL C, PATENTED.

This product possesses the valuable property of producing blacks when diazotized and combined with beta naphthol, in exactly the same manner as reds are produced from diazotizing paranitraniline C. It is therefore very valuable for the production of black and red shades upon a bottom of beta naphthol, when used in conjunction with paranitraniline C. It is especially useful in printing, as the padded goods can be

printed red and black with the two above diazotized compounds. The blacks thus produced are exceedingly fast to light steaming and washing.

To diazotize it, 26.5 parts are dissolved in 125 parts of hot water and 18.2 parts of muriatic acid 21° Beaumé, then cooled with 200 ice and 4.5 parts of nitrite of soda dissolved in 15 parts of water slowly added. The best method of making the printing paste is to add to the above 300 parts of thickening M, and before use add 6.5 parts of acetate of sodium, dissolved in 20 parts of water. The thickening M is 100 parts of wheatflour, 225 parts of water, 150 parts of tragacanth thickening (165:1,000) and 30 parts of acetic acid, 11° Twaddle, boiled together.

AMIDO NAPHTHOL BD, PATENTED.

This product produces blacks when treated with diazotized paranitraniline C, in the same manner as reds are produced from beta-naphthol It is very valuable in producing red and black effects in printing in conjunction with beta-naphthol. The goods are printed with beta-naphthol for red and with amido naphthol BD for black, and the goods thus printed are passed through a solution of diazotized paranitraniline C. The shades are exceedingly fast to light, steaming and washing.

AMIDO NAPHTHOL 3B, PATENTED.

This is similar in its properties and method of application to amido naphthol BD, the shades of black being more of a blue tone than those of the BD brand.

RED DEVELOPER C, PATENTED.

This product possesses the properties of beta-naphthol, when combined with diazotized paranitraniline C, except that the shade is of a much bluer tone. Shades resembling Turkey red can be produced by adding about 10 per cent. of red developer C to the beta naphthol for padding. The fastness of the red thus produced is not lessened in any degree.

PARANITRANILINE RED.

Make up a bath of 1 per cent. of paranitraniline, 5.5 per cent. of nitrite of soda, 2.5 per cent. of hydrochloric acid, 21° Beaumé, and 2 per cent. of acetate of soda. Enter the goods, wash, dry, enter into a solution of beta-naphthol. While para-

nitraniline gives a red or rose according to the concentration, sometimes even a brown, meta-nitraniline will give an orange, dianisidin blues and violets.

In order to obtain a bright color with paranitraniline red it is necessary that the naphthol-prepared cloth should retain its original white look before it goes into the diazotizing bath, otherwise a dull and uninteresting color will result. Betanaphthol absorbs oxygen from the air and becomes brown. To avoid this defect, Lauber and Caberti patented the idea of adding antimony salt to the naphthol solution, keeping the antimony in the solution by the addition of glycerine. The antimony salt is fluoride of antimony and potassium chloride double salt. Knecht recommends the use of antimony hydrate; A. G. Green states that one part of tartar emetic to each four parts of beta-naphthol is effectual. The latter authority also states that equal weights of glycerine and naphthol effectually prevent the development of the brown color.

Inasmuch as oils add greatly to the brilliancy of the color obtained Turkey red oil has for a long time been employed; Messrs. Meister, Lucius & Brüning have now introduced para acid (ricinoleic acid), para soap PN (ricinoleate of ammonia), and para soap RN (ricinoleate of soda), as substitutes for Turkey-red oil.

It will be just as well to mention the different forms in which paranitraniline red is sold. Paranitraniline $C_0H_4 {\stackrel{\textstyle NO_2}{\stackrel{\textstyle \sim}{N}H_2}}$ or its azo derivative are sold in several different forms for the production of azo red upon the fiber. These are:

First, Paranitraniline (free base). Second, Paranitraniline hydrochloride. Third, Paranitrodiazobenzene chloride. Fourth, Paranitrophenyl-nitrosamine.

No. 1 requires the addition of hydrochloric acid and nitrous acid; the second of nitrous acid alone; the third of alkali; the fourth of acid.

Since it has been proved that paranitraline red can be successfully dyed without the employment of ice, its application has very rapidly extended. To get along without the low temperature for which ice has been employed, the usual sodium acetate solution must not be added to the diazotized solution until the latter is required for use.

PARANITRANILIN RED DISCHARGE.

Caustic soda is said to lift the red, and that the complex produced can be best removed by water-glass solution.

If glycerin, glycerides, chlorhydrine, or similar substances be added to tin salts, the discharge can be readily effected.

The ager must be free from oxygen; hence, in presence of aniline black as part of the design, the goods are allowed to hang overnight in an oxidation chamber at 36° to 40° Centigrade before steaming. After steaming, the goods are immediately passed through a tepid bath (40 grms, of strong hydrochloric acid per liter) to remove the different decomposition products formed during the process. Then they are well washed to remove all traces of tin.

When there is a white discharge only, the goods may with advantage be passed through an acidified solution of bleaching powder. This still further improves the white, and the red not only does not suffer, but is brightened by the chlorination. In lieu of a white effect, a colored ground can be obtained by adding to the white discharge (Azorongeant PN of Koechl) gallocyanin, gallamin blue, etc.

PARANITRANILINE C.

Dissolve 10½ pounds of paranitraniline C in 4½ gallons of hot water and 2½ gallons of muriatic acid, 32° Twaddle. When completely dissolved, add, while briskly stirring, 14½ gallons of water. Let the resulting paste cool off completely overnight, then add all at once, and, under brisk stirring, a solution of 5½ pounds of nitrite of soda, dissolved in cold water. Dissolve 11 pounds of beta-naphthol together with 11 pounds of soda-lye, 75° Twaddle, and 11 pounds of soda in about 11 gallons of water, add 6½ pounds of Turkey-red oil, and the base (according to shade required, a very large excess of carbonate of lime or barytes), as well as a sufficient quantity of cold water. Then, while briskly stirring, let the above paranitraniline solution run in. The precipitate forms at once: it is then washed twice with a great deal of water.

RENDERING DIAZO COMPOUNDS STABLE.

This can be done in many cases by mixing with such matter as sulphuric acid, alum, or Glauber's salts.

ALPHA-NAPHTHOL IN BETA-NAPHTHOL.

Alpha-naphthol in the beta-naphthol exerts a detrimental effect in the coloring of the paranitranilin red. In order to prove the amount of alpha-naphthol in beta-naphthol, 0.144 gr. is dissolved in a graduated cylinder in 5 cc. of pure alcohol, and 15 cc. of toluol is added; 0.14 gr. of paranitranilin red is dissolved in 9 cc. of dilute hydrochloric acid, and after this is cold it is diazotized with 1 cc. of normal sodium nitrate solution and then added to the naphthol solution. This is well shaken up and water is added, the two layers which are formed are divided by a separatory funnel, and the color of the toluol solution is compared with a solution which is made in the same way from beta-naphthol containing a known amount of alpha-naphthol. It is possible in this way to determine 0.01 gr. of alpha-naphthol.

REACTION TO DISTINGUISH BETWEEN ALPHA-NAPHTHOL AND BETA-NAPHTHOL.

Cold, saturated, aqueous solutions of the naphthols, made by trituration in a mortar, will give a distinctive coloration with sodium hypobromite. This latter is obtained by mixing 30 cc. of caustic soda of 36° Beaumé with 100 cc. of water and shaking with 5 cc. of bromine. To 10 cc. of the naphthol solution add 2 drops of the hypobromite, when alpha-naphthol will turn to a dirty violet coloration or precipitate, while beta naphthol gives a yellow.

Cross Dyeing.

Wool mordanted with bichromate, woven up with wool which has not been mordanted, and dyed with logwood, the mordanted wool colors black, while the unmordanted is but slightly tinged. If the dyed fabric is topped with a violet dye, as methyl violet, the cloth will have the appearance of a black and violet weave.

The alizarines are ready for this sort of work, for the reason that, if properly done, after dyeing with alizarine, the bath is so thoroughly exhausted that it can be used for dyeing the second color direct; for instance, a blue-red or a yellow-green weave is obtained by dyeing first with alizarine blue, subsequently with some ponceau or bordeau.

A similar effect can be also secured in cotton. A cotton yarn mordanted with tannin and antimony, and woven up with unmordanted yarn and then dyed with a basic color, as methylene blue, and after dyeing washed and then dyed again with a substantive color, as benzopurpurine, a cross color effect is established which is very attractive. Alizarine red gives a bordeaux upon chrome, a bright red with alum, and lilac with iron. Yarn treated with these mordants woven up with unmordanted yarn and then dyed in alizarine gives a quadruple-colored fabric.

Again, in the instances of unions, the wool may be dyed with acid fuchsin, azofuchsin G, scarlet 3R, azocochenille, orange GT, acid violet 4B extra, 5B, cyanol, cyanine Victoria blue, and indigo extract. Use 4 per cent. of sulphuric acid, with 5 to 10 per cent. of Glauber's salts. If the cotton becomes colored much, enough sulphuric acid has not been used. Or the cotton may be mordanted with sumac for three or three and one-half hours. Fuchsin, safranin, auramin, methylene blue, and basic colors are applicable. If Diamine Colors are employed, a full amount of Glauber's salts must be used. To insure an alkaline bath, soca can be placed in the bath. Goods should be entered cold, and worked up very slowly in temperature. If toward the end a dve is needed to color wool and not cotton-supposing that the wool has not taken up as much color as the cotton-then use orange extra, crystal OR, azo-red, formyl violet S 4B, acid green extra concentrated B, alizarine black, or naphthol blue-black.

The Diamine Colors may be divided into:

- (a) Those which dye the cotton darker than the woole.g., diamine violet N, diamine sky blue, diamine new blue,
 diamine black BH, diamine oranges G and D, diamine fast
 yellow A, diamine brown V, etc.
- (b) Those which dye the wool darker than the cotton—e.g., diamine brown B, diamine gold, diamine bronze G, diamine blue 3 R, diamine scarlet, etc.

(c) Those dyeing the cotton and wool to the same shade—e.g., diamine dark blue B, diamine brown 3G, diamine brown N, union black S, diamine red 5B, diamine fast red F, diamine Bordeaux, diamine green.

Wool and cotton can be dyed different shades in one bath. Colors must be used which will dye cotton in acid-bath without staining the wool, as well as wool dyes which will not dye the cotton. Azo red A, acid green extra concentrated, crystal scarlet OR, brilliant orseille, formyl violet, naphthol black, orange crystals, naphthol yellow, and cyanole extra will do the latter work. Diamine Colors which dye both fibers in an acid-bath are used. Diamine brilliant blue, diamine sky blue; then, if for instance, acid scarlet be mixed with diamine blue, a brown and blue shot effect is produced.

COTTON BLEACHING BY ELECTROLYSIS.

A process has been recently devised, and perfectly satisfactory results are said to have been achieved. Cotton yarn is packed in liquors, prepared in a vessel known as the electrolyzer. Through this a solution of rock-salt flows, and at the same time an electric current is passed through the liquid. This bleaching liquid is kept in closed vessels until required. Although the first expenditure is large, the cost of working is said to be low. Two men should bleach and wash off 1,500 pounds of yarn a day.

Applications of Electricity.

Dr. Goeppelsroeder, in Mulhouse, developed ten years ago, and in the Manchester Exhibition of 1889 showed a booth full of samples of dyeings produced by electricity.

It now appears that the Society of Chemical Industry in Basle produces colors by electrizing certain compounds. Furthermore, certain yellows are now obtained by the Badische Anilin- and Soda-Fabrik.

Chlorine obtained by electricity from salt is now being marketed by Arnold, Peck & Co., of New York and Providence, and seems to be of excellent grade. The best bleach will run 35.6 per cent. of chlorine.

The chloride of lime which is now being manufactured in this country gives excellent tests for chlorine strength; the Niagara brand runs 35.6. Two imported bleaching powders gave 35.3 and 35.6 chlorine strength. These figures given are cach the average of three separate determinations. French bleach gave a chlorine strength of 31.47 per cent.

LIQUID SULPHUROUS ACID.

When liquid sulphurous acid can be obtained at three cents a pound, there is small excuse for manufacturers who fail to get a good white.

SIZINGS.

Water-proof: Saturate with gelatine, to which 2 per cent. chrome has been added. Formaldehyde is useful.

Jute bags are sized with milk of lime, and then a mixture of 3 parts oil and 1 part of paraffine.

Another bagging sizing can be made with acetate of lime and Iceland moss.

OIL-STAINS.

The Société Industrielle de Mulhouse has given a medal to the inventor of a solution for the removal of oil-stains from cloth. The composition is as follows:

Water50	parts.
Soap 3.6	44
Ammonia or ammonia soap 0.9	part.
Soda 1.8	44
Oil, such as paraffin-oil, toluidin,	
anilin, and phenol 0.5	66

THE QUESTION OF LIME-STAINS.

The question of lime-stains in goods is of perennial interest, and a dyer has to be eternally vigilant to avoid trouble. A sure way of introducing lime into goods is to have a deficient souring; unless the goods from the lime boil are passed through an acid of sufficient strength, using about two degrees Beaumé, depending upon the weight of the goods, lime-stains are sure to be in the fiber. One of our large bleacheries found out that although the sours were two degrees Beaumé, the acid strength as analyzed in the laboratory was only 0.3 of a degree. The specific gravity was raised by the soluble calcium salts, and the reading by the glass was wholly misleading. Bleacheries should have test solutions to determine the true strength of their acids, chloride of lime, etc. These are readily made for the acids by simply making alkali of proper strength, and adding to it an indicator such as methyl orange; this alkali should be made such a strength that equal volumes of the test solution will exactly neutralize equal volumes of the acid to be tested.

PRESERVATION OF PEROXIDE OF HYDROGEN.

By addition of 2 per cent. of alcohol or ether, solutions of peroxide can be kept for several months. The Société Industrielle de Mulhouse has given a bronze medal to the inventor of this idea.

A bronze medal was also given by the Société Industrielle de Mulhouse for the discovery that naphthaline preserved the strength of solutions of peroxide of hydrogen. The inventor found that with one grain of naphthaline a solution retained its strength almost indefinitely, even at a temperature of 40° Centigrade.

Calcined magnesia will prevent a too rapid decomposition of the peroxide of hydrogen, according to authority.

BLEACHING WOOL WITH PEROXIDE OF SODIUM.

Put into a wooden tank supplied with lead-pipe connection, water enough to cover the wool to be bleached; add 3 per cent. of oil of vitriol, and add peroxide of sodium slowly and with constant stirring, until the liquor is reduced to exact neutrality. It must be remembered that heat will cause a loss of oxygen and must be avoided. Now introduce the wool and make the solution slightly alkaline with some ammonia. Turn on steam and keep bath for one and one-half to two hours at about 140° F. Draw off, wash first with water containing ½ per cent. of oil of vitriol, wash again with water, and dry.

Peroxide of Sodium would appear to be specially adapted to the bleaching of wool fabrics, silks, linen, etc., which suffer more or less by the treatment with sulphurous acid or chloride of lime. But above all it seems to be the best bleach yet employed for mixed cotton, wool, or silk goods in which the animal fiber is positively damaged by contact

with the chloride or the vegatable fiber by treatment with sulphurous acid.

THE WEIGHTING OF SILK.

The weighting of silk has been carried to a degree which would seem as if it would affect the conscience of the most hardened. It is known that tin chloride produces, with tannic acids, particularly catechu and the fibroin of the silk, an insoluble compound. The amount of the foreign matter taken up can be increased, if the silk is previously treated with iron sulphate and ferrocyanide of potash. To be sure, silks treated in this way are only adapted for the dyeing of black, but for the other colors " pink salts." SnCl, 2NH,Cl. can be used. Furthermore, sodium phosphate, followed by water-glass, is applied, and in this way the fiber is enlarged as well as increased in weight. It is not possible to estimate the adulteration either by the touch, or eye, or taste: chemical means are the only methods. Percentage of silk is obtained by making a determination with nitrogen, using the Kjeldahl method. Normal silk boiled, and then dried at the ordinary temperature, holds 15 per cent, of nitrogen. Artificial silk can further be proved by its lack of the characteristic odor of silk when is burns.

DEVING-ROOM DEVICES.

The man in charge of the can-room of a bleachery frequently notices the trouble that the boys have at the dry end of a set of cans, when, under certain circumstances, the dried cloth is being plaited down: it seems that something like an electrical condition of the goods is developed, and the goods wind round the two rollers used in the plaiting.

This trouble can be entirely obviated by a simple device, which should be brought to the attention of the managers of the bleacheries. It consists in a board about four inches wide. held immediately under the roller around which the cloth was apt to wind itself.

Another idea we noted was a device placed at the end of the cans where more than one width of the goods is run at the same time. It was a slight rail-attachment, similar to a fence, going very nearly as high as the plaiting rollers, and preventing the two bundles of dried cloth from running into one another.

ANTIBENZINPYRIN.

Spontaneous combustion frequently occurs in silk works when the silk is worked in benzin. It may be prevented by the addition of antibenzinpyrin, which is the subject of German patent No. 83,048, covering the addition of certain fatty acids to the benzin bath. A 1-10 or 1-100 of 1 per cent, of oleate of magnesia is said to prevent electric sparks. Other fatty acids effect the same result.

EMBOSSING WITH ALUMINIUM.

Extremely pretty effects are obtained in a certain works abroad, by embossing cloth, and then where the cloth has been raised by the embossing to print powdered aluminum. The aluminum is used in an ammoniacal solution of caseine, and to it may be added shellac in borax solution.

ANILINE BLACK.

This black still remains the finest and best black that can be produced by the dyers' art. Properly dyed, it gives the cotton a soft feel and does not rub off. Sansone, in his "Dyeing and Calico Printing," speaks of a one-dip aniline black: "The only difficulty of this otherwise very useful black is the rubbing, for it rubs very badly." He gives tests for aniline black; one being a test with soda and soap along with a swatch of white cotton; the second being a treatment with 10 per cent, sulphuric acid; the third, sulphuric acid; fourth, stannous chlorid and muriatic acid. There are two disadvantages in the production of aniline black; one is the liability of the color to turn green, and the other is the tendency of a tendering of the fiber. The oxidation cannot be stopped at the exact second when the color is developed, and as a consequence the oxidation attacks the fiber. Professor Noelting, in conjunction with Dr. Lehne, published a book on aniline black, in substance reprints of articles by these gentlemen. Another book has recently been published by Dr. Riehmeyer. Among the substitutes proposed are bottoming with Matheson's diamine black, and subsequently topping with weak aniline black. The Farben-Fabriken are recommending aniline chloride as an aniline salt, which does not attack the fiber so much.

The firm of William J. Matheson & Co. bring forward a method of continuous dyeing, Diaminogene B on mercerized cotton goods to replace aniline black. The essential advantages claimed are equality in rank with aniline black as regards shade, fullness, and absolute immunity from tender goods; cost not greater than oxidized aniline black. This method of continuous dyeing is essential in the adoption of machine dveing. The large feeding tank containing the dvestuff solution is placed on a level higher than the dveing-vats. After the dyeing has been completed the goods are washed and then diazotized. Three vats are arranged at such a level that the liquors can be drawn from them. One vat is to contain nitrite of soda and the other muriatic acid; these two articles are run into the first box of the dyeing machine, and the diazotizing is done there. The next box in the machine is to be used for rinsing, and this is done by the use of acidulated water. From the third vat the liquor known as the developing liquor is fed into the third tank. Phenylene diamine, previously dissolved in soda ash, can be used as a developer, also phenylene diamine and resorcin, which combination is usually employed to obtain a shade nearer aniline black. Phenylene diamine and beta-naphthol can be used for blue-black shades.

MERCERIZING.

With the furore over mercerizing cotton into silk, it has now been tried on wool. There has been a good deal of nonsense about the raising of cotton to the grade and rank of silk, because, in the first place, you must have a very good staple, and, after all the labor is put out, is really the gain so very material?

Figures are now to hand showing that while caustic of a specific gravity up to 20° Beaumé reduces the tensile strength very materially, on the other hand, from 20° Beaumé upward, to 36° Beaumé, the strength is less altered, and from 36° on to 50°, a positive increase in tensile strength is secured. What temperature is used in such tests would be very relevant matter, and what contraction is superinduced would be pertinent.

Before passing an opinion on the value of goods said to

be mercerized, it is very advisable to test by washing and drying. Some of the best-looking "mercerized" cloth will lose every particle of its beautiful finish on washing. A pseudo-mercerizing is on the market.

A second application of mercerizing is the very attractive relief effect which can be obtained by printing on to the calico a reserve and then passing the cloth through caustic soda.

A test for mercerized cloth is given in the "Journal of the Society of Chemical Industry," of 1898, on page 43: "Mercerized cotton gives blue color with iodine solution and sulphuric acid." "When mercerized cotton is treated with ammoniacal copper oxide solution, a slight swelling takes place, the central canal usually widens, and the cell wall shows longitudinal striations; but the fiber does not become swollen and constricted in the manner of ordinary cotton, nor are portions of the cuticle dissolved away."

CHROME LACTIC-ACID MORDANT.

The first usage of chrome dates from about 1819, when chromate of lead was used for yellow. In 1832, Camille Koechlin used chrome for the first time as a mordant. Since that time chrome has become an indispensable factor to the dyer and the calico printer. Camille Koechlin, sixty years ago, said, "Chrome has become for our industries the metal of coloration in exactly the same way as iron became for the world the metal of civilization."

The use of bichromate of soda is daily increasing in the practical field, and the more expensive bichromate of potash is being crowded out.

Whether or not lactic acid is better than cream of tartar or sulphuric acid depends considerably on the dye which is to be used. Dyes which alter with chromic acid, such as gallein, alizarine blue, and coeruleine, work well if sulphuric acid is employed in conjunction with lactic.

Good results have been obtained with the following recipe:

Two per cent. of bichromate of potash and soda.

Three per cent. of lactic acid of fifty per cent.

One per cent. of sulphuric acid.

Enter goods at 70° Centigrade, work for half an hour, then boil for an hour.

A good way is to mordant first in bichromate and sulphuric acid and then, after the lapse of half an hour, to add lactic acid.

Chrome has always been considered the mordant for wool and not for cotton. It is now proposed to pass cotton through a solution of bichromate of either potash or soda, and then to develop the hydrate of chrome by applying sulphurous acid. In the presence of excess of ammonia, bisulphite will not react, but so soon as the ammonia is driven off by heating, then the precipitation of the hydroxide of chrome occurs. It is suggested that 50 grams of potassium bichromate be used with 260 grams of bisulphite solution of 56° Twaddle and 100 grams of ammonia. This is raised to a liter, the cotton placed in it for awhile, then dried, steamed, and washed. Considerable chrome is in this way precipitated on the fiber, and alizarine dyes can be used upon the mordant.

E. Knecht suggests that neutral ammonium sulphate should be used to fix chrome oxide on the fibre. In the cold this does not react upon the chromate, but upon steaming it is reduced to chrome exide.

LACTIC ACID.

Patent No. 584,707 covers a method for the manufacture of lactic acid, consisting of boiling under pressure with the milk of lime at a temperature of 130° C., when the carbohydrates, such as sugars, syrups, starch, etc., are altered by hydrolysis, and the calcium lactate is formed. method to make lactic acid is by fermentation.

THE AMEND PROCESS.

The fastest wool colors are produced on chrome mordanted wool, but most all the modes of proceeding agree in one point, namely, that the mordant has to be boiled.

It was a new departure when O. P. Amend suggested mordanting below the boiling-point. The saving of steam will appeal as first advantage of this process to everyone, but by looking more thoroughly other merits will be discovered which will surpass the economy of saving steam.

The boiling chrome liquor attacks the wool. Sometimes the loss in weight of the wool fiber is considerable. It has to be borne in mind that this loss means more than a mere mechanical reduction, but it also depreciates the value of the material. The Amend chrome mordant avoids this, not only saves in weight, but renders the mordanted material of higher quality.

In boiling chrome-mordanted wool the chrome is fixed upon fiber which has been attacked by the chrome liquor. The chrome may be combined with the wool substance and oxidation products. But, according to Amend's mordanting process, chromic acid is taken up by the wool fiber without oxidation. No oxidation products being present, the chrome must be in combination with the wool fiber in the natural state. After the reduction of the chromic acid, Amend's mordant represents a combination of wool fiber and chrome oxide only.

The following is the process: For each hundred pounds of woolen material take six pounds of acetic acid (commercial, 30 per cent.), enter at 140° F. and give thirty minutes. Add one pound of chromic acid in water, let stand thirty minutes, and add to the same bath six pounds of bisulphite of soda and give thirty minutes at 140° F. No wash after this bath is necessary if goods can drain off well.

The chromic acid which is taken up by the wool fiber is reduced to chrome oxide by means of the bisulphite of soda. The appearance of the mordant differs from the usual boiling mordant in not being yellow.

Many of the alizarine colors have been recommended to be used in the presence of ammonia in the dye bath, partly for the reason to make them take somewhat better on the fiber. Ammonia with the Amend mordant works well. There is no chance of any chromic acid being taken off the fiber and the presence of ammonia does not prevent the coloring matter from combining with the mordant.

With the Amend mordant only ½ per cent. of ammonia is used in the dye bath with the coloring matter, and no salts are required. This is another step toward turning out the dyed material in good condition. We commend the Amend mordant to the kind attention it deserves.

MORDANTING AND DYEING WOOL IN ONE BATH.

It has been suggested that the mordanting of wool can be done by using 2 per cent, of bichromate of potash or soda, 3.5 per cent, of lactic acid 70 per cent., and 1 per cent, of sulphuric acid, calculating on the weight of the wool; the wool is to be boiled with bichromate and lactic acid until no more chromic acid can be found, except on the fiber. The dveing can immediately follow, and thus the mordanting and dyeing is done in one bath. For example, a deep blue can be obtained as follows: Mordant one hour and a half by boiling with bichromate, lactic acid, and sulphuric acid in the proportions given above, allow to cool by adding water to 80° Centigrade. Then add 8 per cent. say of Gallein A, paste, and about 12 per cent. of logwood; boil one hour and a half.

CARBONIZING.

The underlying principle is old and well-known, but there have been modifications recently introduced which, in certain specific instances, may be of great value. The prevalent method is to use mineral acids, such as muriatic or sulphuric, or both. Ordinarily, 2 per cent, sulphuric acid is used and the temperature employed is between 70° Centigrade (155° F.) and 72° Centigrade (161.5° F.). The following table is given to show the limits:

Temperature of the hot-room.	Amount of acid used for a two hours' stay in the hot-room.	Amount of acid used for a half-hour's stay in the hor-room.
80° C. (177° F.)	11/2	3 to 7
110° C. (230° F.)	1 to 3	1/2 to 41/2
150° C. (302° F.)	1/2 to 1	1 to 1/2

While the stronger acids have a tendency to yellow the goods, this is not any special disadvantage where the goods are to be dyed. The best authorities counsel a bath of soda, followed by washing. It is suggested, in lieu of the mineral acids, muriatic and sulphuric, that aluminum chloride of 6° to 7° Beaumé be used, the goods allowed to lie some hours, then dry at 50° C. (122° F.), and later 145° C. (293° F.). Magnesium chloride, of a strength of 5° to 6° Beaumé, has been proposed.

PART VII.—STANDARD AND NEW METHODS.

TESTING OF HEMATINE.

The Industrielle Société of Mulhouse is examining a method for testing hematine. The apparatus depends upon optical methods, and uses an instrument somewhat similar to a polariscope. A ray of light traverses the liquid, and by a micrometer screw the depth of color can be adjusted to a standard, and the reading gives the value of the hematine.

LOGWOOD TESTING.

A type should be adopted by mills, whether it be simple logwood chip or logwood extract. The testing of colors should be done in at least three different strengths. It very often happens that a difference not evident on a medium dyeing becomes very glaring on a weak dyeing.

One test on extracts is absolute alcohol. Wash the residue with water, then use the microscope. Chestnut extract and tannins are difficult to isolate in the laboratory, and the dyeing tests frequently come so close together that it takes a long series of systematic experiments to form a final opinion.

A test for chestnut extract is the following: Add tartar emetic, and if there is no precipitate the logwood was free from chestnut extract. Chestnut extract will deposit a gelatinous substance.

As a test for tannin the following has been recommended: Acid bismuth nitrate, with pure logwood extract, will give a cloudiness which redissolves in the excess of the re-agent. Whereas, if any tannin were present, the cloudiness will not disappear, but will precipitate to a marked degree.

Albumen precipitates tannin; acetate of iron can also be

used to precipitate tannin, as well as certain pectic matters and the hematine.

Albert Scheurer proposes a method for the estimation of hematine. The process depends on the fact that aluminum hydrate precipitates completely all the hematine. First, solutions are made of alum and of ammonium carbonate, each to contain 20 grams per liter. Forty cubic centimeters of each of these two solutions are mixed, and after a short boil filtered through a tared filter. This is the blank experiment; simultaneously an identical experiment is made, but 10 cubic centimeters of the logwood extract solution, containing 10 grams per liter, are added. The difference between the blank experiment and the experiment made with logwood gives directly the weight of hematine in 0.1 gram of logwood extract.

At the April meeting of the Société Industrielle de Mulhouse, a method for the analysis of logwood is given, consisting of three dyeings-one upon chrome wool to decide on the hematine; the second, using oxidizing mordants to demonstate the amount of hematoxylin; and the third a dyeing on unmordanted cotton to show the degree of oxidation and the tannins.

SUMACS, ADULTERATION OF AND METHODS FOR THEIR DETECTION.

The Journal of the Society of Chemical Industry has an abstract of an article on the determination of sumac, which is of value to dyers and colorists. Sumac can be adulterated by the leaves of other plants, and this can be detected by a nitrogen determination, an ash determination, or an analysis of the ash. Typical analyses are given, showing the variations between the different methods. Other methods are suggested. One of them is to warm with excess of potash and add a few drops of ammonia molybdate; different colors are obtained, and practice enables the analyst to decide whether there has been adulteration or not. Another method is as follows: A solution of 0.15 gr. of safranine in 1 l. of water serves as the standard of comparison. Five gr. of material are boiled for half an hour with 1/2 l. of water, the liquid cooled, made up to 1/2 l., and filtered. To 25 c.c. of the filtrate in a beaker 5 gr. of a solution of basic lead acetate (specific gravity at $15^{\circ} = 1.184$) and 15 e.c. of caustic potash (specific gravity at $15^{\circ} = 1.155$) are added, and the whole concentrated to 15 c.c. The solution should now be reddish-brown and almost clear. If a noticeable precipitate has formed, or if the liquid is only yellow, the sumac is adulterated. The 15 c.c. are diluted to 250 c.c., and, if necessary, filtered; a solution prepared from pure sumac will not show the tint of the safranine solution.

TESTING OF SUMAC.

ADAPTED TO COLORISTS.

Certain solutions must be prepared, and one of these is the solution of pure tannic acid made to contain one-tenth per cent. tannin; also a solution of indigo extract, about 3 per cent. of indigo paste in water.

First, it is necessary to ascertain the relative value of these two solutions; in order to do this, 20 c.c. of the indigo are put into a white porcelain dish with about one liter of distilled water with a few drops of sulphuric acid; then the permanganate of potash solution is run into this indigo extract solution until the liquid turns pale yellow. This operation is now repeated with 10 or 20 c.c. of the pure tannic acid as prepared above; upon deducting the c.c. obtained in the first experiment from the c.c. obtained in the second, the quantity necessary for the oxidation of the 20 c.c. of the solution of tannic acid is obtained.

The relation of the permanganate to the indigo solution should be such that about an equal number of c.c. of each is required, that is to say, the process works to its best advantage when about 20 c.c. of indigo extract require 20 c.c. of permanganate, and when 10 c.c. of the permanganate of potash requires 10 c.c. of the tannic acid as above described.

About five grams of the sumae extract under examination is weighed out, dissolved in pure water, and raised to one liter. Then 20 c.c. of the indigo extract solution is poured into the white porcelain dish along with a few drops of strong sulphuric acid, with 30 c.c. of the solution of sumac extract, and the whole is raised to one liter. Then the permanganate of potash solution is run in from a burette until decolorization

is effected. For example, 10 c.c. of the solution of permanganate of potash corresponds to 20 milligrams of pure tannic acid; 20 c.c. of the indigo solution are decolorized by 20 c.c. of permanganate of potash solution. Five grams of sumac extract have been dissolved, as it has been said, in one liter of water; 30 c.c. of this solution with 20 c.c. of indigo has required, we will say, 40 c.c. of permanganate of potash to decolorize it; therefore, as 20 c.c. of indigo solution are decolorized by 20 c.c. of permanganate of potash, the remaining 20 c.c. (40 less 20) of the decolorizing solution has been necessary for the oxidation of the tannic acid in the sumac extract.

THE ESTIMATION OF TANNIC ACID.

ADAPTED FROM THE UNITED STATES METHOD.

First the hide powder is tested, and this is done by shaking 10 gm. with about 200 c.c. of water for five minutes. This is filtered through muslin, squeezed out by hand pressure, and then the operation is repeated twice more with the same quantity of water; the last filtrate is passed through paper until a perfectly clear liquid is obtained, when 100 c.c. of the final filtrate is evaporated in a weighted dish; this is dried in a water-jacketed oven until the weight is constant. If the residue should amount to more than 1/10 of 1 per cent., the sample of hide powder should be rejected. Another method of testing the hide powder is to use gallotannic acid, and the hide powder must absorb 95 per cent. of the total solids.

After this preliminary testing of the hide powder the sample under examination is tested for moisture. This is done by using three grams, if it be an extract, and two grams if it is a bark. A flat-bottom dish is used, not less than three inches in diameter.

The second determination is on total solids. If the substance under examination is bark or similar matter, use such an amount of material as to give about 1 gm. of total solids per 100 c.c. of solution, extracting in a soxhlet apparatus; if the substance under examination is an extract, use 20 gm., and dissolve this in 900 c.c. of water at 80° Centigrade, allow it to stand for twelve hours, and then raise to 1,000 c.c. We

are now ready to proceed with the determination of the total solids. This is done by shaking well, and then taking about 100 c.c. in a pipette, using care that the solution should be at 20° Centigrade. Now evaporate in a weighed dish, and dry to constant weight in a water-jacketed oven.

The next step in the analysis is to proceed for the estimation of soluble solids. To do this, 125 c.c. of the solution at 20° Centigrade is filtered through a double-folded filter, returning the filtrate through the filter once. Then evaporate 100 c.c. to constant weight and weigh.

Then comes the determination of tannin substances, and this is done in the following way: Prepare 20 gm. of hide powder by shaking for five minutes with 250 c.c. of water and straining through linen. Repeat the operation three times. Finally, remove as much water as possible by squeezing in a press. Transfer the pressed hide powder to a covered dish, and weigh. Dry a portion approximately equal to one-fourth of the whole to constant weight at 100°. Add the remainder of the hide powder to 200 c.c. of the original unfiltered solution and shake for ten minutes. through linen, collect the filtrate, add 5 gm. of kaolin free from soluble salts, stir well, and filter, returning the first 25 c.c. Evaporate 100 c.c. as before. The weight of the residue must be corrected for the dilution caused by the water contained in the hide powder. The difference between the soluble solids and the last residue obtained is the amount of tanning material absorbed by the hide powder.

TESTING THE NON-TANNIN FILTRATE.

- (a) For Tannin.—Test a small portion of the clear non-tannin filtrate with a few drops of a one per cent. solution of gelatin. A cloudiness indicates the presence of tannin, in which case the determination must be repeated, using 25 gm. of hide powder instead of 20 gm.
- (b) For Soluble Hide.—To a small portion of the clear nontannin filtrate add a few drops of the original solution, previously filtered to remove reds. A cloudiness indicates the presence of soluble hide, due to incomplete washing of the hide powder. In this case, repeat the determination with perfectly washed hide powder.

STRENGTH OF REAGENTS.

Potassium Carbonate. K₂CO₃.—Solution in five parts of water.

Sodium Carbonate. Na₂CO₂.—A solution of one part of the crystallized salt (Na₂CO₂.10H₂O) in five parts of water.

Ammonium Carbonate. (NH₄)₂CO₂.—One part of the crystallized acid carbonate (NH₄)₄H₂(CO₂)₃, in four parts of water, with addition of one part of ammonium hydrate.

Ammonium Sulphide. Normal, (NH₄)₂S.—Super-sulphide, (NH₄)₂S₂. Acid sulphide, NH₄HS.

Sodium Phosphate. Na₂HPO,.—One part of the crystallized salt, Na₂HPO,.12H₂O, in ten parts of water.

Ammonium Oxalate. (NH₄)₂C₂O₄.—One part of the crystallized salt. (NH₄)₂C₂O₄.H₂O, with 24 parts of water.

Potassium Bichromate. K₂Cr₂O₅ or K₂CrO₄, CrO₄.—Solution in ten parts of water.

Potassium Bromide. KBr.—One part dissolved in fifteen parts of water.

Potassium Iodide. KI.-Solution in 20 parts of water.

Potassium Ferrocyanide. K,FeCy,.—The crystallized salt (K,FeCy,.3H,0) dissolved in 12 parts of water.

Potassium Ferricyanide. K,Fe,Cy,...-Solution in 12 parts of water.

Potassium Sulphocyanide. KCyS.—Solution in 12 parts of water.

Ammonium Chloride. NH,Cl.—Solution in 8 parts of water. Magnesium Sulphate. MgSO,.—Solution of the crystallized salt (MgSO, 7H,O) in 10 parts of water.

Calcium Chloride. CaCl.—Solution of the crystals (CaCl. 6H.O) in eight parts of water.

Calcium Sulphate. CaSO₄.—Saturated solution, containing one part of the salt to about 400 of water.

Barium Chloride. BaCl.—Solution of the crystallized salt (BaCl., 2H.O) in 10 parts of water.

Ferrous Sulphate. FeSO,.—Solution of the crystallized salt. FeSO,.7H,O. in 10 parts of water.

Ferric Chloride. Fe₂Cl₄.—One part of the solid salt, Fe₂Cl₄. 6H₂O, to 15 parts of water.

Copper Sulphate.-CuSO,.5H,O.-To 8 parts of water.

Lead Acetate. $Pb(C_2H_3O_2)_2$.—One part of $Pb(C_2H_3O_2)_2$.3 H_2O dissolved in 10 parts of water.

Silver Nitrate. AgNO3.-Solution in 20 parts of water.

Mercuric Chloride. HgCl₂.—One part in 16 parts of water. Mercurous Nitrate. Hg₂(NO₃)₂.—One part dissolved in 20 parts of water acidulated with one part of nitric acid.

A SHORT SYSTEM OF ANALYSIS ADAPTED TO COL-ORISTS AND CHEMISTS.

PRELIMINARY TESTS ON LIQUIDS.

Litmus, curcuma, KI and Pb papers. Try one portion with H.SO, dil., and note the gas evolved; also HCl, dil.; and to this latter add BaCl, for sulphates. In presence of Ag, Pb and Hg, use BaNO,. Try another portion with AgNO, and HNO, for chlorides. Corroborate presence of chlorides by PbA, giving white precipitate soluble in hot H₂O. Iodides will give a vellow. Further, chlorides heated with a mix of K2Cr2O2 and H2SO4 evolve fumes of Cr2Cl2O2. Traces of free HCl can be detected by heating with MnO2 and passing evolved Cl through KI and starch solution. 2KI + 2Cl = 2KCl + I2. FeSO, concentrated solution, with H2SO4 conc. for nitrates; indigo solution also. Magnesia mixture or molybdic solution with HNO, for phosphates. CaCl2 solution gives white precipitate insoluble in CH3COOH with oxalates; a white precipitate soluble in KOH with tartaric acid; and a white precipitate soluble in HCl with H.SO, or H.SO.,

Evaporate a portion to dryness, and follow the preliminary tests for solids.

PRELIMINARY TESTS ON SOLIDS.

Flame Reactions. Beads of PhS. and Na₂B₄O₅.—Heat in a straight and open tube, held obliquely for S; then heat in a bulb-tube, first alone and then with KIISO₄, and afterward with HCl dil.; and then with H₂SO₄ conc. Heat on C alone, and then with NaHCO₅, using Ag coin for S; heat also on C. with KNO₅. KCy, Bi flux. Try solubility in H₂O, HCl dil.; HNO₅; H₂SO₄. Or fuse with K₂CO₅ and then dissolve.

Silver Group.—Add HCl dil. Filter and wash precipitate, which may be AgCl, PbCl₂ or Hg₂Cl₂; wash twice with cold

H₂O and once with hot; save only hot washings; to hot washings add H₂SO₄; white precipitate shows Pb; confirm by K₂CrO₄. To precipitate left on filter add AmOH; if it turns black it shows Hg. Filter and add HNO₈.

Copper Group and Arsenic Group.—Warm the solution and add H₂S; precipitate may be PbS, HgS, Bi₂S₂, CdS, CuS, and As₂S₂, Sb₂S₂, Sb₂S₂, SnS, and SnS₂. Wash the precipitate, and digest with yellow Am₂S, using K₂S in presence of CuS. Note: Am₂S is colorless, and Am₂Sx is yellow; saturate three parts of AmOH with H₂S to make AmSH, and then add two parts AmOH to make Am₂S. Filter off the sub-group and boil off the H.S.

Copper Group.—Boil with HNO₃ and filter; black residue shows Hg. To filtrate, add H₂SO₄, dil., and a white precipitate shows Pb. Filter and add AmOH, when a white precipitate proves Bi(OH)₃ and a blue indicates Cu; filter and add H₂S, and a yellow shows Cd, and a brown precipitate Cu. If brown, solve in KCN and add H₂S, and a yellow precipitates proves Cd.

Arsenic Group.—The preliminary tests are: Heat on C with six times its weight Na₂CO₃ mixed with KCN for the As odor; Sb is volatile and Sn gives a non-volatile coat, going to bluishgreen with CoN₂O₆. Sn in Na₂B₄O₇ with CuO gives a red glass bead. After the H₂S has been boiled off, re-precipitate the sulphides with HCl dil., and then dissolve them in HCl conc. Filter. The residue is As₂S₅, which is taken up by aqua regia, and corroborated by Marsh's test, where it dissolves in NaOCl or Cl water. The liquid holding SbCl₃ and SnCl₄ is evaporated to drive off the major part of the HCl; then add Zn, and in some cases KClO₃ and Pt, when Sb will produce a black stain on the Pt, soluble in HNO₅, dil. For proof of the Sn add HgCl₂ to the solution, which will grow darker and give a white precipitate, showing Sn.

 $SnCl_2 + 2HgCl_2 = SnCl_4 + 2HgCl.$ $2HgCl + SnCl_5 = SnCl_4 + 2Hg.$

Iron Group.—First perform a preliminary experiment to prove presence or absence of oxalates, phosphates, and chrome.

For Oxalates.—One part of the precipitate formed by Am₂S is boiled with a concentrated solution of Na₂CO₂: the filtrate, while still hot, is acidified by acetic acid, and CaCl₂ is added: CaC₂O₄ shows original oxalates. If oxalates were present, dissolve in HCl. and evaporate to dryness, and heat to redness to change all oxalates to carbonates.

For Phosphates.—Use Am₂MoO₄ dissolved in AmOH, and, when clear, pour on to HNO₃ cone. until a permanent precipitate begins to form. From Valentin, 286. For preliminary phosphate test, use the precipitate formed by Am₂S, dissolving it in HNO₃ dil. but hot, filter, and add the Mo solution.

The preliminary for chrome is made by the PhS bead; also by fusing with KNO₃ and Na₂CO₃, which forms K₂CrO₄; filter, neutralize with acetic acid, add AgNO₃, and this gives a brownish-red precipitate, AgCrO₄.

Full Course.-After the elimination of oxalates the preeipitate from Am.S is dissolved in HCl and evaporated several times to dryness with HNO3, or else the residue obtained by the transformation of oxalates into carbonates is dissolved in HNO3 cone. This nitrie-acid solution is heated on a waterbath for half an hour with excess of tin. Then take up with H₂O acidified with HNO₃ and filter; the residue is (PO₄)₄Sn₂ and SnO₂, and the filtrate holds nitrates of Fe, Al, Cr, Zn, Mn. Warm gently the solution holding the iron group, add AmCl, AmOII, and AmoS. This gives FeS, Al2(OH)3, Cr(OH)3 ZnS, MnS, NiS, and CoS. Filter, wash with water containing a little Am-S and finally with water alone. Treat the precipitate with HCl dil. and cold, and then filter; this leaves behind the NiS and CoS. Filtrate is to be boiled with KClO3 erystals until the odor of Cl persists, then nearly neutralize with Na₂CO₂ or NaOH, and allow to stand until it becomes perfectly cold; then add BaCO3 and let stand for some time, shaking occasionally; finally filter; the filtrate (a) holds ZnCl2 and MnCl2; the residue (b) holds Fe₂(OH)₆, Al₂(OH)₆, Cr₂(OH)₆ and the excess of BaCO3. Dissolve in H2SO4 dil., boil, and filter; if necessary, boil and re-filter; to filtrate, add KOH in excess, boil three minutes, and filter; (aa) precipitate is Fe(OII)3 and Cr(OH)3; (bb) filtrate is Al(ONa)3. Fuse precipitate aa with KNO3 and Na₂CO₃, dissolve in H₂O, filter, and add CII₃COOII; Pb(CH₃CO₂)₂, and a yellow precipitate proves Cr. Its solution in HCl shows Fe by the blue with KCy; bb filtrate, with HCl and AmOH giving a white precipitate shows Al₂(OH)₆. Fuse with CoNO, for making eobaltous aluminate.

The filtrate (a), holding the Zn and Mn, is treated with KOH in excess and filtered; (i) to the filtrate add H₂S for ZnS; (ii) precipitate is to be fused with KNO₃ and Na₂CO₃ to

get the green sodic manganate, Na₂MnO₄. Corroborative test for Mn is the violet borax bead while hot, turning to amethyst on cooling. Reducing flame makes it colorless.

The nickel and cobalt come so rarely into color-work and the discrimination between them is so little called for that it hardly seems worth while to give methods in detail. In a borax bead the Ni gives a brown bead and the Co a bluc. Ni with Co will give the blue, and is to be dissolved in aqua regia; evaporate off excess of acid, add KNO₂, and stand; yellow precipitate shows Co; filter, and to filtrate add KOH, when a green precipitate shows Ni. Valentin, 33.

Alternate.—Dissolve in KClO₃ with HCl, evaporate to dryness, dissolve in H₂O, add KCN in excess, 5 per cent. solution, then a drop of acetic acid, and boil a minute; add NaClO in excess and boil again; black precipitate shows Ni; filter off Ni, and test for Co by evaporating to dryness and using Na-B₄O₇ bead.

The borax bead will only show the blue after long treatment with the R. F., in cases where there is much Fe, Cu, Mn, or Ni.

Barium Group.-Add HCl, boil, and filter to remove S: then make alkaline by AmOH, warm, and add Am₂CO₈, and filter. Wash precipitate with hot H.O. dissolve in a little HCl. and add CaSO4. Immediate precipitate shows Ba; precipitate after awhile indicates Sr. To another portion of the precipitate dissolved in the HCl, add H2SO4 and boil to remove Ba and Sr; filter neut, by AmOH and add ammonium oxalate for Ca. Or evaporate the HCl solution with excess HCl, and warm with absolute alcohol and separate to (a) residue and (b) filtrate. (a) If BaCl₂, outer flame in burner is made green, the solution in H₂O treated with K₂CrO₄ gives BaCrO₄. (b) This holds the CaCl₂ and SrCl₂. Evaporate to dryness with HNO₃ several times, and warm with absolute alcohol and filter. (aa) Residue is SrCl₂ (flame test), and the filtrate (bb) is Ca-(NO₂), which colors the flame vellowish red, and with oxalate ammonium gives CaC2O4.

Am, K, Na, Mg Group.—Treat original liquid with NaOH and heat for Am; AmOH, and Na₂HPO₄ for MgHPO₄; also blp. and CoN₂O₆. Flame tests for Na and K. Special for K is that PtCl₄ gives 2KCl.PtCl₄ which ignites to Pt and KCl; ammonium gives 2AmClPtCl₄ which ignites to Pt only. Tartaric acid

with K and Am gives similar precipitates, but the K ignites to alkaline residue.

ARGOL ANALYSIS.

1. METHOD FOR ESTIMATING THE TARTRATES IN AN ARGOL.

Into a 100 c.c. flask put 7.52 gm. of the argol to be tested, and add 30 e.c. HCl (320 gm, of acid to the liter). After shaking thoroughly, let it stand overnight. Then make up to 100.5 or 101 c.c., according to the amount of insoluble matter in the argol. Next heat 25 c.c. of a solution of potassic carbonate (of 328 gm, to the liter), and slowly add to it 50 c.c. of the filtered argol solution. Having boiled this solution five minutes, wash it into a 100 c.c. flask and raise to that volume. Then filter half of the solution and evaporate it to 15 c.c. Add 3 e.c. of glacial acetic acid and from 100 to 150 e.c. of 95 per cent, alcohol, taking about five minutes to add the latter. Allow the mixture to stand half an hour, so that there will be complete formation of crystals, and then decant and wash the crystals on a filter with alcohol till they are free from acetic acid. When the crystals are well drained return them to the dish with the filter-paper and titrate with deci normal solution of sodic hydrate, using phenol phthalein as indicator.

2. METHOD FOR TESTING THE POTASSIC BITARTRATE IN AN ARGOL.

Boil 3.76 gm. of the argol five minutes with 750 c.c. of water, cool solution, and make it up to a liter. Filter 500 c.c. of the solution through a dry filter-paper, evaporate this filtered solution on a water-bath to dryness, and after the residue is dry let it stand on the bath half an hour. Moisten the residue with 5 c.c. of water. When the mixture is cold, add to it 100 c.c. of 95 per cent. alcohol, stir well, and let it stand for half an hour. Put the liquor and crystals on a dry filter-paper, and when the crystals are well drained return them to the dish with the filter-paper and add 100 c.c. of water. Heat the mixture and titrate with deci normal solution of sodic hydrate, using phenol phthalein as an indicator.

ZINC DUST ANALYSIS.

A new method has been proposed, based on the reduction of ferric sulphate, and the amount of ferrous salt formed is determined volumetrically with standard permanganate solution. Of the zinc dust in question, 1/2 gr. is put into a stoppered 250 c.c. flask, together with 25 cc. of water. Shake until the zinc is well suspended, and then introduce 7 gr. of chemically pure dry ferric sulphate.

After shaking for fifteen minutes, the reduction will be complete; then add 25 c.c. of strong sulphuric acid, fill up with water, mix, and allow to settle. Draw off 20 c.c. of the liquid, add the same volume of water, and titrate with permanganate. The amount of ferrous salt is immediately indicated, and the percentage of metallic zinc may be calculated from the formula:

 $Fe_2(SO_4)_3 + Zn = ZnSO_4 + 2FeSO_4$

SOAPS AND SOAP ANALYSIS.

ADAPTED TO THE NEEDS OF COLORISTS.

Caustic soda permits of a fictitious "body," and the soap has a tendency to hold a larger percentage of water without giving itself away so readily.

What the management of a mill desires to know is the moisture, the kind of oil or fat employed, and whether there is any free caustic, free carbonate, starch, insoluble matters, silicate of soda, Glauber's salts, or unsaponified fat in the soap.

To obtain this the analysis proceeds in the following way: 5 to 10 gm. of fine shavings are dried at 60° to 70° Centigrade, and later between 100° to 105° Centigrade to constant weight. Soaps containing free caustic should be dried in carbonic acid, free air, or the water is determined by difference after all the other constituents have been estimated.

The second test is the determination of unsaponifiable and uncombined fat. This is done by taking the soap which was dried in making the moisture determination, and extracting with ether; this extract is then evaporated and weighed, and the percentage of unsaponified fat is thus obtained.

The third determination is for free caustic. The residue from the fat determination is dissolved in alcohol, filtered, and to the filtrate phenolphthalein is added; if the color becomes pink it indicates free caustic alkali, but if there was no change of color there was no free caustic. In the event of the soap turning to pink, decinormal or centinormal acid is run in until the color returns yellow. The reading obtained gives a basis for calculating the amount in percentages.

The fourth determination is for any free carbonate. To make this determination, the insoluble residue from the previous test is used. It is dissolved in warm water and is titrated with decinormal HCl acid, methyl orange being used as an indicator.

Total Alkali.—The total alkali consists of the free caustic and free carbonate and the combined alkali. Thirty grams of soap dissolved in hot water with 50 c.c. of normal sulphuric acid. Filter the whole or an aliquot part of the same. The filtrate is titrated back with caustic soda, methyl orange being preferably used as indicator. The amount of alkali combined with the fatty acids is obtained by subtracting the sum of the free caustic and carbonated alkali from the total.

The fifth determination is for insoluble matter. Any insolubles will remain on the filter from the last test. The filter-paper is dried and the estimation made by simple calculation into per cents. This insoluble residue may consist of water-soluble salts, generally chloride, sulphate, and carbonate of the alkalies, water-glass, borax, etc., or it may consist of water-insoluble mineral substances like chalk, clay, silica; or it may contain organic substances, as starch and glue.

Alkali, carbonated and borated, are determined by extracting with cold water and determining in aliquot portions of carbonate, silicate, and borate of alkali.

Sodium-chloride and sulphate are determined in an aliquot part of water extract by means of silver nitrate and barium chloride. The part insoluble in water is ignited to destroy organic matter, weighed, and then washed and analyzed qualitatively and quantitatively. In the portion insoluble in alcohol, water extracts dextrine, and this can be precipitated from the water solution by alcohol.

The presence of starch in the insoluble residue is discov-

ered by the microscope, and the reaction with iodine solution. The starch can be determined by converting into sugar by means of dilute sulphuric acid, neutralizing the solution with barium carbonate, filtrating, and determining with Fehling's solution. To test the insoluble portion for glue, it is extracted with warm water, the solution gelatinizes on cooling, and gives a precipitate with tannin solution.

The sixth test is the test for combined fatty acids. The alcohol filtrate obtained from the determination made for free caustic is largely diluted with water; enough acid is added to separate the fats. These fats are taken out with ether, weighed, and reported as fatty acid.

Determination of Total Fat.—The fatty acids separated from the soap by the following processes should only be considered as such, when no fat. resin, or unsaponifiable matter is present. In order to determine the amount of fatty acids combined with the alkali, the amount of the foregoing impurities must first be determined and subtracted.

Five to twenty grams of soap dissolved in a small quantity of water, shaken up in a separatory funnel with an excess of diluted sulphuric acid or hydrochloric acid. The separated fatty acids are dissolved with petroleum ether. Petroleum ether solution is then evaporated in a tared flask, and residue weighed. In expressing the results of analysis, the amount of fatty acid has to be calculated to fatty anhydrides.

The question as to whether free fat acids are present may sometimes be settled by Jacobsen's method of adding a little rosaniline to the oil. If free fatty acids are present, the oil turns red in color in consequence of the formation of rosaniline oleate. More important is the adulteration with resin and with hydrocarbon oils. In the absence of free fatty acids, resin may be isolated from fixed oils by agitating the sample with moderately strong alcohol, separating the spirituous solution, and evaporating it to dryness.

The separation of the resin acids from free fatty acids is best effected by a method based upon the ready solubility of silver resinate in ether, and the almost complete insolubility of silver oleate, etc., in the same menstruum, even in the presence of a small quantity of alcohol. The exact separation of the mixed fatty acids is hardly possible. Perhaps the most satisfactory method known is based on the fractional pre-

cipitation of the alcoholic solution of the acids with magnesium acetate. This salt precipitates acids of the stearic series more easily than it does oleic acid and its homologues, and, of the different homologues of the stearic series, those of the highest molecular weights are thrown down first.

The Koettsdorfer number, or the "saponification equivalent," is often a key to the character of the oil or fat. It is determined as follows: About 1.5 to 2.5 gm. of the fat is treated with 25 c.c. of one-half normal alcoholic potash; when saponification has taken place, 1 c.c. of an alcoholic solution of phenolphthalein is added, and the liquid titrated with one-half normal hydrochloric acid. A blank experiment is then made by titrating 25 cubic centimeters of the alcoholic potash alone, and the difference in the volumes of the acid used gives the volume of the potash solution neutralized by the fat, each 1 c.c. corresponding to .02805 gm. of potassium hydrate, whence the saturation equivalent is easily calculated. The saponification test is used in connection with many others for the identification of oils, and is known as the saponification number, or Koettsdorfer value or number.

TESTING FOR ACIDS IN GOODS.

This can sometimes be done by dyeing the goods in alkali blue.

PART VIII.—SUNDRIES.

BUYING ON SPECIFICATION.

THE MODERN METHOD OF PURCHASE.

For a number of years the Pennsylvania Railroad has been purchasing by specification. When this plan was first entered upon there was a great deal of trouble, but to-day the very dealers who were the greatest objectors are now the most desirous of having more specifications drawn.

By a specification is meant a specific, definite, concise, and detailed order. Nearly all law suits come through lack of definition in the contract. One man says one thing and means another. In logic also it is a first rule that proper definition is the prime requisite.

To give some suggestions the following might be cited. Individual cases require modification, but these specifications are fairly representative.

TOILET SOAP .- This must show on analysis:

- Not more than one-fourth of one per cent. of foreign mineral matter.
- Not more than one-fourth of one per cent, of carbonated alkali reckoned as carbonate of soda (Na₂CO₃).
- 3. Not more than one-fourth of one per cent, of uncombined caustic alkali reckoned as caustic soda (NaOH).
- Nor should the soap leave an odor of cocoa-nut oil, palm oil, or palm-nut oil on the hands.

COMMON SOAP .- This must not show on analysis:

- More than one-half of one per cent. of foreign mineral matter.
- 2. More than one-half of one per cent. of carbonated alkali reckoned as carbonate of soda (Na $_{\circ}$ CO $_{i}$).
- 3. More than one-fourth of one per cent. of uncombined caustic alkali reckoned as caustic soda (NaOH).

4. Nor must the soap leave an odor of cocoa-nut oil, palm oil, or palm-nut oil, on the hands.

Scouring Soap. The material desired under this specification is a soap which must not contain:

- 1. More than 30 per cent. of water.
- 2. More than 1 per cent. of free carbonate, unless stipulated.
- 3. It must have no free caustic soda.
- 4. It must contain no silicate of soda nor rosin soap, unless stipulated.
 - 5. There must be no foreign mineral matter.
- 6. It must not contain nitro-benzol or other substances to disguise unpleasant odors. The odors should not be disagreeable when rubbed in the palm of the hand.

FULLING SOAP.—The material desired under this specification should meet the following conditions as near as possible:

- 1. It must not contain more than one-fourth of one per cent. of foreign mineral matter.
- 2. It must contain no silicate of soda nor rosin, unless so stipulated.
 - 3. It must have no free caustic soda.
- 4. It must not contain more than one-fourth of one per cent. uncombined carbonate of soda, unless so stipulated.
- 5. It must contain no organic matter foreign to a pure soap.
- 6. It must contain no nitro-benzol or other substances to disguise the odor. The odor should not be in the least disagreeable when rubbed in the palm of the hand.
- 7. The soap must be made up approximately of 7 per cent. combined alkali, 63 per cent. fatty anhydrides, and the balance water.

PROVISIONAL SPECIFICATION FOR GALIPOLI SOAP FOR A SILK MILL.—The material desired under this specification is soap which must not contain:

- 1. More than 30 per cent. of water.
- 2. More than one-tenth of one per cent. free carbonate.
- 3. More than one-fourth of one per cent, of foreign mineral matter.
- It must contain no silicate of soda or resin unless otherwise stipulated.
 - 5. It must contain no free caustic soda.
 - 6. It must contain no nitro-benzol or other substance to

disguise the odor. The odor should not be in the least disagreeable when rubbed in the palm of the hand.

7. The soap must be made up from pure olive oil.

CRYSTALLIZED GLAUBER'S SALTS.—Crystallized Glauber's salts or normal crystallized sulphate of soda (Na₂SO₄ .10 H₂O).

- 1. Must not contain material amounts of iron.
- 2. Must not contain more than 56 per cent. water.

CALCINED OR DESICCATED GLAUBER'S SALTS.—The material it must not contain:

- 1. Material amounts of iron.
- 2. More than 1 per cent. of water.
- 3. It must not be acid.

SAL SODA.-The sample should have 21 per cent. alkaline strength, and approximately ten molecules of water.

PROVISIONAL SPECIFICATION FOR CAUSTIC SODA.-The material desired under this specification is sodic hydrate (NaOH), as free as possible from other substances. It may come in drums, or as a powder. It must not contain notable quantities of:

- 1. Iron.
- 2. Carbonate.
- 3. Water.

SPECIFICATION FOR CAUSTIC POTASH .- The material desired under this specification is potassic hydrate (KOH), as free as possible from other substances. It must not contain notable quantities of:

- 1. Iron.
- 2. Carbonate.
- 3. Water.
- 4. Less than 97 per cent. caustic potash will be returned.

PROVISIONAL SPECIFICATION FOR MURIATIC ACID.—The material desired under this specification is commercial muriatic acid, 20° Beaumé, chemically known as hydrochloric acid, HCl. as free as possible from all other substances.

- 1. A sample containing any considerable amount of iron will be rejected.
- 2. Acid of 20° Beaumé at 15° Centigrade should contain 32 per cent. HCl.
 - 3. It must evaporate clean.

SPECIFICATION FOR SODA ASH is to be drawn in accordance

with the alkaline strength. The relation between the values is given herewith:

Na ₂ O	NaCO3	English Alkali.	Liverpool and New York Alkali Test.
46.5		47.11	48.00
47.0	80.37	47.62	48.50
49.0		49.64	50.58
51.0	87.19	51.67	52.64
53.0	• • • • •	53.70	54.70
55.0		55.72	56.77
55.5	• • • •	56.23	57.29
56.0	95.74	56.74	57.80
56.5		57.24	58.32
57.0	97.45	57.75	58.83
57.5		58.26	59.35
58.0		58.76	59.87
58.5	100.02	59.27	60.38

SPECIFICATIONS FOR SAL SODA.—The sample will be considered good which has 21 per cent. alkaline strength, and has approximately ten molecules of water.

BLEACHING POWDER.—The strength of the article in available chlorine will serve as a basis upon which bills will be paid. Prices should be quoted on 35 per cent. chlorometric strength, and for each per cent. over or under 35 per cent. chlorometric strength 1/35 should be added or subtracted.

NITRITE OF SODA.—The specification calls for sodic nitrite (NaNO₂), as free as possible from all other substances.

The sample should test above 97 per cent, of sodium nitrite. Blue Vitriol.—The material desired is normal crystallized sulphate of copper, as free as possible from other substances.

- 1. It must not contain more than $\frac{1}{4}$ of 1 per cent, crystallized sulphate of iron.
- 1. It must not contain more than $\frac{1}{2}$ of 1 per cent. of impurities.

SAL AMMONIAC.—The material called for by this specification is granulated chloride of ammonium (NH₄Cl), as free as possible from any other substances.

- 1. It must not contain less than 65.15 per cent. of chlorine.
- 2. It must not contain less than 31.2 per cent. of ammonia (NH_4) .

COCOANUT OIL.—Cocoanut oil is desired as free as possible from all other substances.

- 1. The iodine number should be between 8 and 9.
- 2. The saponification number should be above 265.
- 3. The specific gravity at 99° Centigrade, taking water at 15.5° Centigrade, should be above 0.87.
- 4. It should be soluble in two parts of 90 per cent, alcohol at 60° Centigrade.

WHEAT-STARCH.—The material desired under this specification is wheat-starch, free as possible from all other substances.

- 1. It must not contain notable per cents. of ash.
 - 2. It must not contain more than 15 per cent. of water.
- Under the microscope it must show the characteristic granules of wheat-starch as given in photographs of wheat granules in standard works.

CORN-STABCH.—The material desired under this specification is corn-starch, free as possible from all other substances.

- 1. It must not contain notable per cents. of ash.
- 2. It must not contain more than 15 per cent, of water.
- Under the microscope it must show the characteristic granules of corn-starch as given in photographs of corn-starch in standard works.

TEXTILE PRIZES.

The Industrial Society of Mulhouse, Alsace. Germany, has issued the programme of prizes which are to be competed for in 1899. This society annually offers prizes, medals, and honors. A partial list is given herewith.

A medal of honor will be given for a synthesis of the coloring principle found in cochineal; for a study on cochineal carmine: for a red which can be used on tannin mordant, which will be as bright as alizarine red and fast to soap.

A prize is offered for a way to account for the differences in aniline blacks.

A prize is offered for a work of scientific value concerning the transformation of cotton into oxy-cellulose. A paper of merit upon the synthesis of one of the many natural coloringmatters used in the dyeing industry will be honored. A silver medal is offered for a coloring-matter which can replace logwood in its different applications, having the advantages of logwood in point of stability and fastness. A meritorious study in iron mordants and the rôle which these mordants play in dyeing will receive a medal.

A medal will be given for a blue similar to ultramarine blue in fastness and shade, which blue must be applicable to cotton without the aid of albumen or other thickening agent. There is a silver medal for a yellow which can be fixed in the same method as alizarine and will possess the solidity equivalent to alizarine yellow. A medal is offered for a reserve for steam colors particularly adapted to wool and removable by a simple washing in water in the same. A silver medal is offered for a powder which can be printed on the cloth and give the same effect as gold or silver. This powder must resist washing.

A medal of honor, either silver or bronze, according to the merit of the work, will be given for the best practical manual upon the bleaching of cotton, wool, wool with cotton, silk, flax, linen, and other textiles.

 Λ medal is offered for a research upon the preparations and applications of hydrogen peroxide as a bleaching agent of textile fibers.

A silver medal will be given for a marking compound suitable for cotton to be dyed in reds or deep colors. There is a silver medal offered for a practical process which will remove stains from cloth. This new process must be applicable and at a price which will not increase the cost of the goods materially. The removal of oil-stains by hand is not a permissible solution of the problem.

A silver medal is offered for a thesis explaining the cause which makes different kinds of cotton bleach defectively.

DYEING WHITE.

There are two patents recorded by Mr. Carter. One of them speaks about the state of art, phrasing the matter as follows:

"Prior to my invention it has been the custom, in the art of calico printing particularly, to imprint upon the fabric the designs with a pigment consisting of oxide of zinc or sulphate of barium, to which albumen has been added, and to subsequently subject the fabric to the effect of heat, so as to decompose the albumen, whereby the latter will mechanically bind the pigment upon the fabric. It has also been proposed to print upon the fabric, where a white effect is desired, with tungstate of soda to which gum or starch is added, and subsequently passing the fabric through a solution of barium chloride, whereby insoluble tungstate of barium will be produced as the coloring-matter. This color, however, has been more or less fugitive, and it is the object of my invention to increase its permanence,"

This invention consists in subjecting tungstate of soda in such a way that it remains permanent on the fiber. The inventor has discovered that a sulfocarbonate of cellulose, when applied to a fabric and subjected to heat so as to be decomposed, possesses the property of adhering with such tenacity to the fabric as to be practically incapable of removal, and he bases his invention upon this discovery. He adds to a sulfocarbonate of cellulose a suitable white pigment, preferably tungstate of barium, and when the former is subjected to heat it be fixed with almost absolute permanence upon the fabric. The sulfocarbonate of cellulose which is used in carrying out the improved process is known in the market as "viscose."

FREEZING MIXTURES.

Substances to be mixed in parts by weight:	
1.—Water	1
Ammonium nitrate	1
2.—Snow	1
Sodium chloride	1
3.—Snow	1
Calcium chloride	1.3
4.—Snow	1
Sodium chloride	0.4
Ammon. chloride	0.2
5.—Snow	1
Sodium chloride	0.416
Ammon, nitrate	0.416

BRIGHTER COLORS.

It is said that colors can be brightened by using a decoction of soap root (Quillaya Saponaria) in the dye-baths in conjunction with the colors dyeing in neutral baths.

TUNGSTEN AND MOLYBDENUM AS MORDANTS.

A recent number of the Revue Générale des Matières Colorantes has an article on the Mordants of Tungsten and Molybdenum by Edward Knecht. It is only worth the while to give the conclusions, namely, that these mordants do not fix the colors upon the fiber with the same satisfaction which can be obtained by using the mordants from chrome.

CEMENTING CLOTH TO IRON.

A good cement for fastening cloth to iron is said to be made from linseed oil, rendered dry by boiling with litharge mixed with well-dried pipeclay; then thin with oil of turpentine.

AN ADHESIVE CEMENT.

Equal proportions of gum tragacanth and tapioca ground together and heated with water at about 70° F., after which there are added about 50 per cent. of gum and an equal amount of starch, and the whole mixture heated at from 70° to 120° F. The cement thus produced is said to be superior to any hitherto known.

FULLERS' EARTH.

The best is found in Buckinghamshire and Surrey, England. When good it is of greenish-white or greenish gray color, falls into powder in water, appears to melt on the tongue like butter, deposits very little sand when mixed with boiling water, and is soft and greasy to the feel.

DRYING OF LAKES.

There has been brought before the Committee of Chemistry of the Société Industrielle de Mulhouse a new process for the treating of lakes. The process consists of precipitating upon a thin sheet of metal the lake to be treated. This thin sheet of metal passes over into a current of warm air, where the precipitate is dried. The process has been invented by M. Alph. de Huillard of Suresnes.

SHRINKAGE IN FLANNELS.

According to a French patent, shrinkage of flannels can be avoided by passing the goods through a strong solution of borax mixed with albumen, followed by a dilute acid with or without an addition of alum.

HOW TO WASH WOOLENS.

The washing of woolens in the laundry to avoid shrinking is a household problem solved as follows: Soak as hot as can be borne by the hand in water in which soap has been boiled. To about every six gallons of water add from three to five tablespoonfuls of liquid ammonia, which removes grease deposited by perspiration. After an hour's soaking, wash out, and refrain from rubbing or kneading the fabric. To thoroughly remove the soap, rinse out twice in lukewarm water. Iron while still damp, stretching the article to the necessary length and width. The odor of the ammonia disappears when the soap has been removed.

An alternative mode is the cold-water method. In an ordinary bucket of water, dissolve a piece of good soap, and add a teaspoonful of ammonia. Into this put the goods to soak six hours. Express the suds, by drawing through the hand or with a wringer, and riuse in cold water till the water flows off clear and clean.

APPLICATIONS OF AMMONIA.

Ammonia can be used with soap to advantage. Certain manufacturers are using sal ammoniac and speak very highly of its advantages. Where a vat with tightly fitting lid is to be had, it should be employed, since the heat is not only saved, but maintained equably, and there is not the same loss. It cleans out soap and sweetens the goods. Ammonia should not be neglected in the dye nor the finishing rooms.

ARTIFICIAL SILK.

The furore over artificial silk has somewhat subsided, and it is the general impression that too high an estimate was made of the qualities of the product. It is, however, unquestionable, the statements of the many interested detractors to the contrary notwithstanding, that it will be a serious rival to the natural silk in a great many applications. It is said that the prospective uses of artificial silk made under the Lehner patents are daily increasing. The company's works at Glattburg, near Zürich, Switzerland, are said to produce 3 cwt. of yarn each week, and the works are fully occupied day and night.

There are three methods to-day for the manufacture of artificial silk. One is the Chardonette, another is the Vivier, and the third is the Lehner. The Chardonette process forces a collodion solution through capillary tubes, and these filaments are eoagulated, and four of them are twisted into one thread. This is de-nitrated to remove its explosive qualities.

The Vivier method starts with nitrocellulose, fish sounds, and gutta percha.

The Lehner process mixes together the collodion with a solution of pure silk waste.

The most appalling defect in artificial silk is its inability to withstand moisture. Even in the form of cloth, pure artificial silk tears with the most melancholy ease.

It has been discovered that cellulose can be dissolved by heating it with a strong solution of chloride of zine and then adding hydrochloric acid.

A new process for artificial silk consists of three treatments: First, the threads of cotton are passed into a solution of cellulose, through a solution of cupra-ammonia, thio-carbonates, or nitro cellulose; second, the thread is then passed into a bath which coagulates these compounds of cellulose upon the fiber, and water, benzene, toluylene, or xylene are used; third, the process is completed by a bath acting as a fixation; for instance, dilute acid if the Schweitzer solution has been used, an alkaline sulpho-hydrate, an acetate of iron, or an iron chloride if nitro-cellulose has been used. In this way a thread is obtained having the appearance and all of the tinetorial properties of natural silk.

Silk gloss upon cotton yarn, hosiery, etc., is said to be producible by treating the material at 40° Centigrade with acetate of lime 5° Beaumé, then, without rinsing, working at 50° Centigrade with a bath of Marseilles soap, finally upon a 10 per cent. bath of acetic acid and drying without rinsing.

A NEW ALBUMEN PREPARATION.

The Compagnie Parisienne de Coleurs d'Aniline have a patent for treating egg albumen with formaldehyde. The mixture is allowed to stand several days, when the water is added, and the formaldehyde then boiled out. The residue is filtered and dried in a vacuum pan. The product can be precipitated by acids and alcohol, and dissolved in water, and is not precipitated by alkalies.

RAMIE.

California has come forward as an aspirant for favor in the rhea fiber industry. Mr. Felix Fremerey has long been considered an authority on the subject, and his experiments at Bakersfield, Kern County, Cal., have given great impetus to the cultivation of fibrous plants.

HEMP.

California is coming into prominence as a successful manufacturer of hemp, and the yield of water-retted fiber at the farm of Mr. John Heaney, at Gridley, Cal., is about 2,600 pounds to the acre.

FLAX.

Oregon is said to be one of the best flax countries. The Oregon Women's Flax Fiber Association is engaged in bringing flax into prominence, and their efforts are untiring in advertising the possibilities of the fiber. This industry is but two or three years old, and yet the product is meeting with a good demand.

A professor at Lille has discovered the micro-organism indispensable in the retting of flax. It is the "bacille amylobacter." Other micro-organisms develop in the water during the operation, having a very noxious influence and destroying the amylobacter bacillus. The inventors constructed a building in which the decomposition of the fiber, with the aid of the amylobacter bacillus, and its drying, occur simultaneously, and the whole operation lasts five days.

TEXTILE SCHOOLS.

The Lowell Textile School was opened the first of February of this year under the very best of auspices. We give some pictures of some parts of this school and also of the Philadelphia school on pages 157 and 158. The Philadelphia Textile School, which has won for itself a wide reputation during the fourteen years of its existence, and the proposed textile schools at Fall River, Paterson, and Charlotte, give most encouraging guarantee that the next generation of mill managers and superintendents will have a training requisite for the carrying on of their industries in the most enlightened way. We have long looked for a union between practical and theoretical training, and these schools are our guarantee that this will be done and in the best manner.

In Germany there is a well-nigh universal effort to give technical training a prominent place in education, and all sizes of cities have good trade schools (Fachschulen). Year-Book cannot do better than to quote a paragraph written by Principal L. W. Miller. of Philadelphia Textile School. "That masters and employers have reached a stage of development where they are willing to make it a part of their duty to provide systematic instruction in their trades by means of schools which are open to the apprentices of their rivals in business, as well as to their own, shows how much better than their brethren in America they have learned the importance of making common cause against a common enemy. In America we are still to a considerable extent in a state which, in this respect, is representative, not of a social order, but of savagery; that is to say, it is so far from being the rule that our employers are willing to interest themselves in making general the education which would tend to improve the quality of their productions, that the contrary would probably be more nearly true. Too many of them would rather fear than welcome the effect of this education, which simply means that they would prefer to hunt by themselves in a wilderness rather than make those concessions and render that service to the common safety and welfare which civilization demands. This is the most pressing lesson which American industry has to learn, and Germany is the country in which to learn it."

The highest types of schools of this sort in Germany are the Kunst-Gewerbe Schulen of Berlin, Nürnberg, Munich, Crefeld, and Mühlhausen, with which the admirable schools of Switzerland at Luzerne, Zürich, and Geneva are quite worthy of being classed.

In England there are the Manchester Technical School, the Leeds School, and the Central School at South Kensington, London.

The technical instruction committee of the city of Manchester recently presented its report of a deputation appointed to visit the technical schools of Germany and Austria.

"It is not, however," the committee points out, "only in the domains of science that Germany is making great progress. In almost every town visited by the deputation fine industrial art museums were found, arranged with the express purpose of cultivating a knowledge of what has already been accomplished in the production of fine examples of color, design, and workmanship. Every technical school has its special museum of objects applicable to its purposes. Notably was this the case in Berlin, Vienna, Nürnberg, Crefeld, and at Düsseldorf, in which latter place the Industrial Art Museum is said to be the finest in the Rhineland. At Nüremberg there has been recently erected, at a cost of \$250,000, a fine industrial and trade museum, known as the Bavarian Museum, to which it is intended to add laboratories and class-rooms for chemical technology."

TREATMENT OF BOILER SCALE.

Analyses of boiler scale show that it is usually made up in the following way:

·	Per cent.
Sulphate of lime	. 56.5
Carbonate of magnesium	. 19.8
Carbonate of lime	. 18.0
Oxide of iron and aluminum	. 0.7
Silica	. 3.8
Moisture	. 1.6

Carbonic acid holds the lime and magnesium in solution in the form of bicarbonate. Hence, if something is added to take away that excess of carbonic acid, lime and magnesium are precipitated. Where the lime and magnesium salts are in solution as bicarbonates, this simple addition of a substance designed to remove the excess of carbonic acid will cure the water with no further trouble. The reagent to effect this purpose is usually lime-water itself.

Sulphate of lime in the water is not altered by this process, and hence the lime-water cure is useless for such waters. Soda ash should be used. Quite frequently barium chloride can be used as a remedy for water holding sulphate of lime. Then the calcium chloride is subsequently removed by the addition of soda ash. Carbonate of barium is said to be useful where the water holds sulphate of lime, and in this case there are produced two insoluble precipitates, namely, carbonate of calcium and sulphate of barium.

A large number of substances have been suggested, and the basal idea is to form a slimy liquid which will not harden. Sugar, syrups, starch-meals, glue, tannic acid, potatoes, oils, tar, fats, etc., are used in this way. A large number of antiincrustation powders are on the market.

COSTLINESS OF HARD WATER.

A pound of carbonate of lime or magnesia precipitates ten pounds of soap. Therefore, the evil of hard water is somewhat calculable. Not altogether, because there are losses from partly perfect goods concerning which no account can ever be taken. Iron, magnesium, calcium, and aluminum salts can be removed with attention, and then the plague of precipitated soap is avoided.

TAKING SAMPLES OF WATER FOR ANALYSIS.

A new gallon bottle and a new cork should be used. Rinse the bottle thoroughly with some of the water to be analyzed. Then fill the bottle not quite full, taking care not to have the water which goes into the bottle touch the hands. In case the water is from a spigot or cock, the end of the spigot or cock should be cleaned both inside and outside and then the water allowed to run a few minutes before sample is taken. All general information in regard to the source of supply, possible contamination, or any peculiarities of the water should be stated. Write on the label the date and hour when

the sample was taken, and for identification sign it with your name.

Boiler waters have been treated with tannic acid, which produces tannate of lime, and this being slimy will not form boiler scale. The idea of using compounds which would produce slimes and not mineral decompositions is illustrated by many of the boiler compounds at present on the market, particularly starch, potatoes, dextrine, and even glue and albuminous substances, stearine, tar, whale oil, any fatty substances or a soap. Boiler compounds have been analyzed which contained 75 per cent, logwood. Each case stands on its own merits and has to be treated in strict accordance with the individual requirements. An analysis is always necessary. It takes but very little care to obviate very large losses.

DRINKING WATER TEST.

Hager's test for potable water is a tannin solution made up of one part tannin, water 4 parts, alcohol 1 part. The addition of this solution to potable water should cause no turbidity, even after standing some time. Another test with water is to bottle some of it and keep it corked for a few days. If it is malodorous on opening, it is a suspicious case, unless it be a natural sulphur water. Another rough and ready test is to evaporate to dryness and observe the residue.

Still another experiment which may give suggestions concerning the nature of a water is to fill a good sized white glass jar, and allow it to stand for a day or so. A clouding will give a hint of bicarbonates. The natural sediment will give hints to an experienced eye.

The total solids in water supplied to American cities where a fair degree of purity is maintained, expressed in parts per million, are: Boston, 47: Philadelphia, 134: New York, \$2: Washington, 165: Brooklyn, 64: New Orleans, 340: Indianapolis, 290, and Minneapolis, 156.

More than five grains per gallon of chlorine (71.4 parts per million), accompanied by more than .08 parts per million of free ammonia and more than .10 parts per million of albuminoid ammonia, is a clear indication that the water is contaminated with sewage, decaying animal matter, urine, etc., and should be condemned. Eight-hundredths parts per

million of free ammonia and one-tenth part per million of albuminoid ammonia render a water very suspicious, even without much chlorine.

Albuminoid ammonia, over .15 part per million, ought to absolutely condemn a water which contains it. solids found in the water should not exceed 40 grains per gallon (571.4 parts per million).

METHOD FOR ANALYZING FATTY OILS.

· Take specific gravity and melting point. Use ether to find insolubles.

Five-ten grs. of the sample, previously melted is necessary, is filtered through a dry filter, unless quite clear.

RESIDUE may contain salt, curd, water, sand, and insoluble matter. It may be washed with ether, dried and weighed, then ignited and reweighed; difference la organic matter.

THE CLEAR OIL is shaken out in a separator with water and other, or recently distilled carbon bisulphide. The aqueous liquid is separated, and the oil solution again shaken with water, if the previous treatment was found to remove anything,

may contain soap and is evaporated at 100.

AQUEOUS OIL SOLUTION.—Shake again with dilute sulphuric acid and separate. If an aliquot portion of the oil solution leaves no ash on ignition, thus proving the absence of metallic compounds, the shaking ont with water as above, and with sulphuric acid. can be omitted. Wash the residual oil repeatedly with water till the equeous solution gives no test for acid with litmus.

ACID LIQUID may contain sulphate of alnminnm and heavy metals previonsly existing as soups.

SOLUTION OF OIL IN ETHER OR CARBON BISULPHIDE.-Evaporate, saponify, shake repeatedly with petroleum ether.

Petroleum ether solution is evaporated to determine the amount of mmeral

Aqueous sointion is acidified with petroleum ether and further examined to determine the nature of the futty oils.

LUBRICATING OILS.

A good lubricating oil (1) should decrease the friction; (2) it should not lose its lubricating qualities by standing in the air; (3) it should not have any chemical action on the metals with which it comes in contact; (4) it should possess a certain degree of viscosity, so that it is neither pressed out from between the grinding surfaces, nor by quicker movement of the machine thrown off; (5) at the temperature at which it is used it should give off no inflammable gases or vapors. Mineral oils, fatty oils, and mixtures of the two

are used as lubricants, additions of resin oils and drying oils decrease the quality.

The specific gravity and viscosity of the oils (and the cold test, if the oils must withstand the cold of winter) should be determined.

Mineral oils, as a rule, are free from acid. Sulphuric acid remaining from the refining may be detected by shaking with warm water and testing with methyl orange, the water layer should not become red.

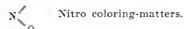
Nitro naphthalin is added to lubricating oils in order to destroy the fluorescence of the mineral constituent. It may be detected by extracting with alcohol. After evaporation, this alcoholic extract will leave long transparent yellowish acicular crystals.

CLASSIFICATION OF ORGANIC DYESTUFFS.

In the Journal of the Society of Dyers and Colourists, S. de Kostanecki classifies organic dyestuffs. He has arranged them according to the nature and the number of the chromophors. and presents the following tabular classification:

A. DYESTUFFS CONTAINING A SINGLE CHROMOPHOR.

- C: C. Dibiphenylene-ethene.
- C: 0. Oxyketones, oxycoumarins, oxyflavones, oxyxanthones.
- C: N. Auramine, thioflavine, quinoline vellow.



- N: N. Azo colors.
- N: N: O. Azoxy colors.

B. DYESTUFFS CONTAINING SEVERAL CHROMOPHORS.

- (a) STREPTOSTATIC CHROMOPHORS (KETONE TYPE).
- C:C) Unsaturated oxyketones, indogenides, oxyindogen-C:C) ides, indigo.
- C:0) Oxydiketones, oxydixanthones. C:05

C:N)
C:N)
Hydrazone dyestuffs.
N:N)
N:N)
Disazo dyestuffs.

(b) CYCLOSTATIC CHROMOPHORS (QUINONE TYPE).

	C:C.	$\mathbf{G}:\mathbf{O}.$	C : N.
C: O	Aurins	Oxyquinones.	
C : N	Basic coloring matters of the triphenyl methane group. Pyronines	Indophenols Nitrosophenols.	Indamiens. Azines. Safranines. Indulines.

(c) STREPTOSTATIC AND CYCLOSTATIC CHROMOPHORS.

This group comprises several complicated coloring-matters, such as alizarin blue, styrogallol, etc.

BALING.

The old-fashioned cubical cotton bale contained about 500 pounds of fiber, but it is now passing out of vogue, and the new cylindrical bale is taking its place. There is now an improvement made in the new bale, namely, that the bale is put up just thirty-five inches wide, in order to adapt it to the opener at the mill.

ANOTHER NEW BALE.

The American Cotton Bale Company is about to place on the market a new style of packing for cotton. Substantially, it is a thin galvanized iron box, held in position by steel straps. The bale weighs 18 pounds. The old style of baling weighs 28 pounds. The first form which the inventor made was an asbestos covering, held in shape by wire; this was too expensive. The second form was similar to the telescope-bag idea. The present form is a very handsome-looking bale weighing 500 pounds, and is making everywhere a very satisfactory impression among the manufacturers. The particular advantage of this bale over the round bale is the fact that

it can be filled by an inexpensive machine. The round bale requires a very expensive machine. The machine for packing this new bale costs from \$125 to \$150.

BASIC COLORS.

M. Horace Koechlin states that nitrosophenol and the naphthols are very strong mordants for the basic colors.

DYES FOR LEATHER.

Black Striker.-A powdered striker for use in logwood dveing, of uniform strength and tone.

New Leather Black.-An aniline black for producing blueblack shades upon leather.

Tannin Brown B .- Adapted equally for the dveing of mordanted cotton and of leather. A clear, pure brown-vellow of good fastness.

THE NEW TARIFF.

The most important acid, sulphuric acid, is taxed 3/c. a peund. Muriatic, nitric, carbolic, and oxalic acids remain duty free. Tannic acid decreases 10c. per pound. Bleaching powder is to pay 1/5 c.: last year there was imported into the United States 103,513,977 pounds, valued at \$1,579,358; this will probably yield a very considerable revenue unless our manufacturers decide to buy in this country. Coal-tar dyes are raised from 25 to 30 per cent.; last year there was imported over \$3,000,000 worth. Aniline salts are free. Also "alizarine, natural or artificial, and dves derived from alizarin or anthracin." Nearly \$1,000,000 worth of alizarine came in last year. Coal-tar, benzol, toluol, and naphthalene remain free of duty. Extracts of logwood alter from 10 per cent. to %c. Logwood extract to the value of over \$1,500,000 was imported last year. Extracts of quebracho alter from 10 per cent. to 1/4c. a pound. Extract of sumae from 10c. to %c. a pound; ground sumae, 3 to 10c.; cutch and gambier are free. Indigo extracts are removed from the free list and charged 3/c. a pound; carmined, 10c. a pound; indigo itself is free; over \$1,673,000 came in last year. Bichromate of potash is to pay 3c. in lieu of 25 per cent.; bichromate of soda, 2c. instead of 25 per cent.

This will further aid in the desirable substitution of the sodium for the potassium salt, because the tariff will increase the present disparity of 2e. a pound between the bichromes. Similar remarks might be made on chlorate of potash, which is to pay $2\frac{1}{2}$ c., and chlorate of soda, 2c. Caustic potash pays 1e.; caustic soda, $\frac{3}{4}$ c.; last year there was imported over \$1,000,000 worth of caustic soda. Soda ash is to pay $\frac{3}{8}$ c. About 120,000 tons are imported into the country.

BLEACHING.

A report from the Norwich Bleaching, Dyeing, and Printing Company states that in the last ten years 570,424,917 yards of cloth have been finished.

INVENTIONS MADE IN THE MILL.

The question concerning the ownership of inventions made by employees arises at times, and it would seem as though the management of mills would follow the method employed at the large establishments abroad, namely, make contracts with the help, so that when anything is invented, using the chemicals of a mill, it becomes the property of the firm. A man anxious for employment will never hesitate to sign such a contract if it is insisted upon at the start. It is quite another thing, however, after a man has been employed for a while and gets a warped idea of his own consequence.

It may readily be a close question whether a chemist or other employee who has done something new has a property right in the novelty, which is a *de jure* right or a *de facto*. If the invention was worked out with the chemicals and machines of the mill, the mill is apt to think that it is the owner in equity. If the workman has used extra time, then he believes himself to be the legal owner in fee.

Industrial confusion which results can be obviated by foresight in contracting against the contingency. The case of Dobson rs. Dempsey; the case of the Silver Spring Co. rs. Woolworth cost much money and trouble, all of which could have been sayed.

COLLODION COVERINGS.

An English patent states a method for giving textiles, such as cotton, linen, hemp, jute, wool, and even leather, wood, gutta percha, and rubber, a color and a silk gloss in one bath. The process passes the articles through an ether-alcohol solution of collodion, to which any suitable dyestuff has been added, and then entering a closed chamber for removing the alcohol and ether and recondensing them.

The fiber is previously treated to a two per cent. solution of starch and then dried. Addition of olive oil to the collodion solution aids in rendering the result less hard. The colors are said to be absolutely fast to light, air, washing, acid, alkali, and chlorine. Whether this process is applicable to selvedges is not yet certain.

PEGAMOID.

This widely and cleverly advertised article seems to be a pliable celluloid. The suppleness is given by the use of castor-oil and the like. The uses to which this pegamoid is applicable are so manifold as to be almost bewildering. The offices of the company display numerous articles "pegamoided," and ranges from an imitation leather to wall-paper and painted articles. The manufacture of a "leather" from cotton cloth, and its application in upholstery reminds one of pantasote, which latter article is used for imitation leather window shades and gives creditable service.

ARTIFICIAL LEATHER.

An English patent covers an artificial leather prepared as follows: Unvulcanized rubber is dissolved in benzine or other solvent. Leather waste is added, and suitable coloring agents may also be employed. This is well mixed and the paste is squeezed out from the box and passes between rolls. The sheet is removed to a drying-shed and is subsequently embossed or treated as desired.

WOOL WASHINGS AS A FERTILIZER.

An analysis of wool washings, made with reference to its approximate value as a fertilizer, gave the following results:

	Per eent.
Moisture at 100° Centigrade	. 41.13
Dry matter	. 58.87
·	

The dry matter left behind contained:

	$^{\rm P}$	er eent.
Potassium oxide		10.15
Phosphoric acid		0.10
Nitrogen		1.09

The commercial value of these ingredients per ton of the original substance amounted approximately to \$12.40. Samples of raw wool tested for potash give the following results:

	Per	cent
Potassium oxide soluble in water	3.	92
Potassium oxide soluble in diluted hydrochloric acid.	4.	.20

The scarcity of a good quality of carbonate of potash for manurial purposes in case of tobacco and similar industrial crops ought to encourage attempts to turn the concentrated potash liquor into a source of revenue.

FINENESS OF WOOL FIBER.

Seventy-five thousand measurements were made under the microscope in the Department of Agriculture. The Department said that it is possible to produce in the United States as fine wools as can be produced in any part of the world.

RESTORING RANCID TALLOW.

It is said that raneid tallow may be returned to usable condition by melting and adding 2 per cent. of chalk, 5 per cent. of lime water, then boiling until a scum no longer rises. This scum should be skimmed off, and the tallow should become of a clear golden yellow color. Five per cent. of sodium chloride is added, the whole mixture boiled five minutes

longer, and then the liquid should be strained through a fine cloth.

CUTCH AND GAMBIER.

Cutch can be distinguished from gambier by treating the sample submitted for examination with normal caustic potash diluted with water four times, and then shaking up with benzole. The gambier will show a green fluorescence.

A NEW SOLVENT.

Formine is an ester formed with formic acid by heating glycerine and oxalic acid to 140° to 190°. It will dissolve many of the hitherto unused aniline dyes which are now insoluble in the ordinary solvents.

STAIN REMOVERS.

Resin stains are dissolved by chlorhydrins, especially if warm. Dichlorhydrin is a product of the action of hydrochloric-acid gas on glycerine. It boils at 176° Centigrade, and therefore is not readily inflammable. It has a pleasant, sweet. ether-like odor. As water decomposes it, the user must take care that the goods are not wet, because the acid produced by its decomposition would ruin certain goods. Since water cannot be used with it, it should be noted that the things which can be used as diluents are benzol, but not benzine, absolute alcohol, chloroform, and carbon tetrachloride.

Epichlorhydrin is made by alkalies on dichlorhydrin. Its boiling point being 117° Centigrade, it is more inflammable than chlorhydrin. It possesses an alcoholic odor, and it is not decomposed by water.

STAINS FROM STEAM.

Anti-crustating compounds, such as petroleum and coal-tar in the boilers, while in many instances are of value in preventing boiler scale, can be very disadvantageous in the matter of producing stains. It is a matter of common knowledge that many oils will distill with steam, and therefore, where vats are heated by use of dry steam, oils of various kinds may be introduced into the vat.

STAINS OF NITRATE OF SILVER.

To remove the stains produced by nitrate of silver, use a concentrated solution of cupric chloride, and then wash with hyposulphite of sodium.

NUMBERING THE GRAPHIC FORMULÆ.

The numbering of the graphic formulæ when the molecule gets somewhat complex is sometimes very confusing. When one ring is used the numbering is as follows:



But when the ring is double, then the numbering and the naming is considerably more complex. The following shows a method of numbering and the names and the combinations:

1:2 Ortho; 1:3 Meta; 1:4 Para; 1:5 Ana; 1:6 Epi; 1:7 Kata; 1:8 Peri; 2:6 Amphi; 2:7 Pros.

MARKING GOODS.

The following things are used at different bleacheries: 'Tar and turpentine, gas-tar alone or mixed with lampblack, printers' ink, copal varnish and black pigments, manganese chloride fixed with alkali, red ochre boiled with cotton-seed oil in the proportions of 750 c.c oil to one kg. of earth.

Aniline black can be used in this way: Make up two solutions, A and B. (a) Boil 170 gr. of starch in 1 liter of water, and, when cold, mix in 60 gr. of chlorate of potash and 48 gr. of copper sulphide. (b) Boil 170 gr. of starch in 1 liter of water as before, and, when cold, stir in 170 gr. of aniline salts.

Use equal amounts of each. Copper sulphide made by heating with continual stirring, 250 gr. of sulphur, 1 liter of caustic soda of 36° Beaumé, and 1,200 gr. of copper sulphate in 20 liters of water. Filter before using.

MISAPPLICATION OF SULPHURIC ACID IN WOOL-DYEING.

No dyeing assistant is more open to misuse than sulphuric acid. The object of using sulphuric acid at all is to set free the color, and as it is relatively quite a cheap article, many dyers use too much of it. If a dyeing bath for woolen cloth is too acid, the dye leaps on to the fiber and irregularities are shown in the result. Certain colors are more sensitive than others to the aid of sulphuric acid, and where a dye is very quick to go on to the fiber, it is often essential to boil the goods in a bath containing only the dyestuff and Glauber's salts, and subsequently adding acid.

ACETIC ACID AS AN AUXILIARY.

Of all acids, acetic acid is perhaps the most useful. There are many dyestuffs on the market which have the capacity to dye too quickly in presence of sulphuric acid. For such dyes acetic acid is well adapted. It causes a slower exhaustion of the dye-bath than sulphuric acid, and as a result the shades obtained are even. It ought to be used for half-woolen goods, since sulphuric acid weakens cotton and takes off the color. Many dyers do not recognize the value of this acid, but acquaintance will no doubt cause a more extensive use of it.

COLOR CONTRASTS.

Colors may be divided into primary, secondary, and tertiary classes:

PRIMARIES .- Red, yellow, and blue.

Secondables.—Green, by mixing blue and yellow. Orange, by mixing red and yellow. Purple, by mixing red and blue.

TERTIARIES.—Russet, by mixing orange and purple. Citrine, by mixing orange and green. Olive, by mixing purple and green.

The following is a tabular view of complementary colors:

Red to green. Yellow to purple. Blue to orange. Green to red. Purple to yellow. Orange to blue.

One of the chief things in all combinations is that the contrast shall be distinct and well balanced.

Colors which are complementary become intensified by contact. If you place together two colors which contain the same common element, such as yellow and green, they suffer; yellow and orange will not give a good combination, because the yellow element is too evident, detracting from the red in the orange composition.

HELP IN CASE OF ACCIDENTS.

Drowning. 1. Loosen clothing. 2. Empty lungs of water by laying body on its stomach, and lifting it by the middle so that the head hangs down. Jerk the body a few times. 3. Pull tongue forward, using handkerchief, or pin with string, if necessary. 4. Imitate motion of respiration by alternately compressing and expanding the lower ribs, about twenty times a minute. Alternately raising and lowering the arms from the sides up above the head will stimulate the action of the lungs. Let it be done gently but persistently. 5. Apply warmth and friction to extremities. 6. By holding tongue forward, closing the nostrils, and pressing the "Adam's apple" back (so as to close entrance to stomach), direct inflation may be tried. Take a deep breath and breathe it forcibly into the mouth of patient, compress the chest to expel the air, and repeat the operation.

Burns and Scalds. Cover with cooking soda and lay wet cloths over it. Whites of eggs and olive oil. Olive oil or linseed oil, plain, or mixed with chalk or whiting. Sweet or olive oil and limewater.

Lightning. Dash cold water over a person struck.

Sunstroke. Loosen clothing. Get patient into shade, and apply ice-cold water to head. Keep head in elevated position.

Mnd Dog or Snake Bite. The cord tight above wound. Such the wound and cauterize with caustic or white hot iron at once, or cut out adjoining parts with a sharp knife. Give stimulants as whiskey, brandy, etc.

Fainting. Place flat on back; allow fresh air, and sprinkle with water. Place head lower then rest of body.

Cinders in the Eye. Roll soft paper up like a lamplighter, and wet the tip to remove, or use a medicine dropper to draw it out. Rub the other eve.

Fire in a Building. Crawl on the floor. The clearest air is the lowest in the room.

Fire from Kerosene. Don't use water, it will spread the flames. Dirt, sand, or flour is the best extinguisher, or smother with woolen rug, table-cloth, or carpet,

Suffocation from Inhaling Illuminating Gas. Get into the fresh air as soon as possible and lie down. Keep warm. Take ammonia-twenty drops to a tumbler of water, at frequent intervals; also, two to four drops tincture of nux vomica every hour or two for five or six hours.

ANTIDOTES FOR POISONS.

Induce vomiting, by drinking hot water or strong mustard and water. Swallow sweet oil or whites of eggs.

SPECIAL POISONS AND ANTIDOTES.

Acids. Muriatic, oxalic, acetic, sulphuric, nitric. Soap-suds, magnesia, lime-water.

Prussic acid. Ammonia in water. Dash water in face. Carbolic acid. Flour and water, mucilinginous drinks.

Alkalies. Such as potash, Ive, hartshorn, ammonia. Vinegar or lemon-juice in water.

Arsenic, rat poison, Paris green. Milk, raw eggs, sweet oil, lime-water, flour and water.

Bug poison, lead, salpetre, corrosive sublimate, sugar of lead, blue vitriol. Whites of eggs or milk in large dones.

Chloroform, chloral, ether. Dash cold water on head and chest. Artificial respiration. Piece of ice in rectum. No chemical antidote.

Carbonate of soda, copperas, cobalt. Soap-suds, and mucilaginous drinks.

Iodine, antimony, tartar emetic. Starch and water. Astringent infusions. Strong tea, tannin.

Mercury and its salts. Whites of eggs, milk, mucilages.

Nitrate of silver, lunar caustic. Salt and water.

Opium, morphine, laudanum, paregoric, soothing powders or syrups. Strong coffee, hot bath. Keep awake and moving at any cost.

Strychnine, tincture of uux vomica. Mustard and water, sulphate of zinc. Absolute quiet. Plug the ears.

POSTAL INFORMATION.

FIRST-CLASS MATTER.

This class includes letters, postal cards, and anything scaled or otherwise closed against inspection, or anything containing writing not allowed as an accompaniment to printed matter under class three.

The rate on matter of this class is two cents per ounce or fraction thereof.

SECOND-CLASS MATTER.

This class includes all newspapers, periodicals, or matter exclusively in print and regularly issued at stated intervals as frequently as four times a year.

The rate on matter of this class is one cent per pound or fraction thereof.

THIRD-CLASS MATTER.

Mail matter of the third class includes printed books, pamphlets, engravings, circulars (in print or by the hectograph, electric pen, or similar process), and other matter wholly in print, proof-sheets, corrected proof-sheets, and manuscript copy accompanying the same.

The rate on matter of this class is one cent for each two ounces or fraction thereof.

FOURTH-CLASS MATTER.

Fourth-class matter is all mailable matter not included in the three preceding classes, which is so prepared for mailing as to be easily withdrawn from the wrapper and examined. It embraces merchandise and samples of every description, and coin or specie.

Such articles as poisons, explosives, or inflammable articles,

live animals, insects, or substances exhaling a bad odor will not be forwarded in any case.

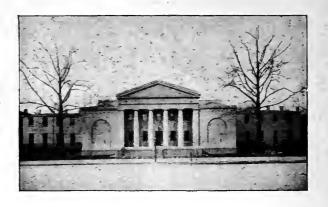
The regulations respecting the mailing of liquids are as follows: Liquids, not ardent, vinous, spirituous, or malt, and not liable to explosion, spontaneous combustion, or ignition by shock or jar, and not inflammable (such as kerosene. naphtha, or turpentine), may be admitted to the mails for transportation within the United States. When contained in glass bottles or vials, such bottles or vials must be strong enough to stand the shock of handling in the mails, and must be enclosed in a wooden or papier-maché block or tube not less than three-sixteenths of an inch in the thinnest part. strong enough to support the weight of mails piled in bags and resist rough handling; and there must be provided, between the bottle and its wooden case, a cushion of corkcrumbs, cotton, felt, asbestos, or some other absorbent, sufficient to protect the glass from shock in handling: the block or tube to be impervious to liquids, including oils, and to be closed by a tightly fitting screw-lid of wood or metal, with a rubber or other pad so adjusted as to make the block or tube water-tight and to prevent the leakage of the contents, in case of breaking of the glass. When enclosed in a tin cylinder, metal case, or tube, such cylinder, case, or tube should have a screw-lid with a rubber or cork cushion inside in order to make the same water-tight, and should be securely fastened in a wooden or papier-maché block (open only at one end), and not less in thickness and strength than above prescribed. It would be well always to consult the postmaster in reference to the proposed mailing of liquids. The limit of admissible liquids and oils is not exceeding four ounces, liquid measure.

Limit of weight of fourth-class matter (excepting liquids), four pounds.

The rate on matter of this class is one cent for each ounce or fraction thereof.



VIEWS IN THE DYEING DEPARTMENT OF THE LOWELL TEXTLE SCHOOL.



The Philadelphia Textile School issues an attractive catalogue and announces a course of study which goes far toward making it unnecessary for American students to go abroad for their training in Textile Chemistry, Dyes and Dyeing. A visit to the school impresses one with the fact that there is the right atmosphere about the place. This right atmosphere, or an earnest and general air of strict attention to the matter in hand, finds its best exemplification in very few American institutions. It is, however, the pre-requisite to the manufacture of well-equipped alumni, which alumni are the measure of success or the non-success of any place of learning.

There may be some question whether the course of study goes far enough, but that is a matter most readily remedied as time goes on.

PART IX.—DYE ANALYSIS.

DYE ANALYSIS.

EXPLANATION OF THE TABLES.

The following is a tabular view which has been used in the laboratory successfully for locating a dye.

The tables are arranged in the spectrum order—violet, blue, green, yellow, orange, and red. The reagents employed are sulphuric acid, water, hydrochloric acid, caustic soda, and a few other reagents of incidental advantage for characteristic reactions.

The method of testing is to place concentrated sulphuric acid in a porcelain dish and then to scatter the dye in question on to the sulphuric acid. The color will either change to one of the spectrum colors or remain unchanged, and the colors to which the dyes turn have been written in columns arranged in spectrum order, and the name of the dye which has been changed is written in that column marked violet, blue, green, etc., to which the acid has altered it.

Then water is added to the test made with concentrated sulphuric acid. There is a change, as a rule, in the color, and the name of the dye is written in the column marked the color to which the dye turns.

A water solution is made of the dye and hydrochloric acid is added; the changes in the colors have been arranged in spectrum order, and, as before, the name of the dye changing respectively to violet, etc., is written in that column.

The water solution is furthermore treated with a caustic soda solution.

Abbreviations used are sol, for solution; ppt. for precipitate. Unless otherwise mentioned, solutions are intended.

The numbers placed after each dye are the numbers from "Schultz's Tabellarische Uebersicht der Künstlichen Organischen Farbstoffe," 3rd edition, Berlin, 1897.

VIOLET.

REACK : Find montred violed R	Rodalsh-gray.	Genetine (Continued, Clematine M. N.	
Violate : Tordomy velor Compare velor Hordomy O O V		Heliotrope B, 9 B Transin Heliotrope Cilradi Ameriysis violed Howeinn Howeinn	
Indian : Chromotrope 8 B 100.		Mauvein	
Braux: Olivomotope 10 B Faak violet reddidt Dimah violet N Congo corfutt (4	Green blue. Phrty green-blue. Green-blue.	White posts Olivaine vlote, Manye dye Affiliation vlote.	
Heliotrope 2 H. 185. Oxamin yielet 188. Azonimyo H. 188.	Clear blue.		def. Olive-green
Congo cortinta B. 211. Holiotrope B. 222. Azo violet. 243. Hoseina viote. 243.		Phenanduo Phenanduo Pripartue Tyvaline	
Oallo eyanine DH, BS	Cornilower blue,	Lydlin, Purpor	
Prune pure	Corntlower blue,	HROWN: Chrome violet pasto	Yellow-brown,
Gracen : Bart vlotet blubdi	Dirty green.	Red violet 5 R exten	Yellow-brown.
Violet neutro extra		Reginn purple	
Methylene ylatet 3 R A		Obronic violet up	Yallow-brown.

RED-Continued. Pist ncid violet A 2 R.	Hilack: First violet reddish	First violet reddish. 145. On dilution red-violet	212. 214. 219.	Methyl violet B and 2 B. Methyl violet V 3. Violet de Paris. Violet an Methyl B. Violet an Methyl B. Violet an Methyl B. Violet activities and the blue then Violet. Crystal violet 5 B O, O. Violet C. Zathyl violet. Zathyl violet.
Brown-yellow,	Brown.yellow.	Brown-yellow, Orange-yellow, Orange-yellow, Bed-yellow,	-366. Red-yellow.	Scarlet-red. Blue-red. Yellow-red. Oringe-red.
305,	305.	31. (313. (314. (317. (318.	386.	36. 8 38. 1 38. 1 319.
Fallow Hoffman's violet Hown-yellow Hoffman's violet B and z B Methy! violet V il. Yelet A bris. Methy! violet V il. Yelet A bris. Yelet wolldow Hown Yelet William Yelet William Yelet William Yelet William Yelet Yelet William Yelet Yelet	Crystal violet Crystal violet Violet C Acthyl violet Henzyl violet Faris violet B T B Methyl violet B T B Methyl violet B T B Methyl violet B T B Violet B B T T B Violet B B T T B	Red violet 5 R S. 311. Acid violet 4 In N, 6 B, 7 B, N 312. Red violet 4 R S. Red violet 4 R S. Red violet 5 R S. Red violet 5 R S. Red violet 5 R B. Red violet 6 R S. Red violet 6 R S. Red violet 7 R S. Red violet 8 R S. R S.	Gallein in paste. Gallein A in paste. Alizarine violet. Anthracene violet.	Wool violet S 36. Scarlet-red. Wool violet A B S Scarlet-red. Victoria violet S B S S S Blue-red. Akad violet B S S Blue-red. Akad violet B S S S S S S S S S

VIOLET—Continued.

First blue.	Olive - green, then	Olive, then green, then blue. Olive-green, then	Violet sol., later blue flocks.	Blue, then violet. Blue, then blue-violet, then red-violet.	462. Blue, then violet-red.	Blue, then red-violet.	Blue, then red.	Blue, then blue-violet. First blue, then red.	First green, then blue, then red-violet.
Violer—Continued. Paraphenylene violet473. Flrst hine.	BLUE: Azo violet	Acid violet 4 B N, 6 B, 7 B, N312 Ulive, then green, Formyl violet S 4 B	Fast acid blue B	Violet neutral violet B. Fast neutral violet B. Violet neutra solide B. 460 {Blue, then blue-violet, Violet neutro solide B.	Methylene violet R R A	Clematine	Heliotrope B, 2 B	Amethyst violet	Mauveïne Rosolan. Violet paste Chrome violet. Mauve Mauve dye Anileïne.
	306 Xellow - green, then F green - blue, then violet.	319. First violet.red, then blue-violet.	348. Vine flocks. 448. First blue, then violet	460. Silve, then blue-viol. let, then red-violet, First blue, then red-violet.	465. First blue, then blue- violet.		(First oreen then	467. blue, then red-vio-	
VIOLET—Continued. Benzyl violet press		Violet fut arctiff 1935. Acid violet 6 B N	Violamine B. Salva (blue flocks.) Neutral violet	Fast neutral violet B	Amethyst violet	Violet paste. Chrome violet	Manye dye. Anileine. Aniline violet.		Indisine Phenamine Phenamine Phenameine Phenameine Tyraline Tyrlan purple

First green, then blue, then red-violed.	Red-brown.	f First dark brown, then blue-green. Lemon yellow. Orange-red.	Yellow-red ppt. Fuchsine red. Puchsine red. Yiolet-red, then bineviolet.
GREEN—Continued. Murveine Roadin Roa	Tyraline. Tyrino purple Lydino Baown: Chrome violet paste	YEL	
First green, then blue, then red-vio-let.	302. { Olive - green, the on green, then blue. 303. { Manue - green, then blue - green, then violet. 304. { Green, then blue luto } 304. { Violet.	Green, blue, then violet. Yellow - green, then green - blue, then violet.	Olive, then groon, then blue. b. but brown, then blue-green. First olive-green, then green, then blue. dirty green plt.
Antiline viole. Antiline viole. Antiline viole. Violene Perkina violet. Perkinanine Phonamenine Phonamenine Pyraline Tyraline	Hofmann's violet 302 (Olive - green, the on Methyl violet B and 2 B 302 (green, then blue. Methyl violet V 3 303 (green, then blue. Violet an Méthyl B 303 (violet wolling (inon plus ultrn). 303 (violet kolunie (inon plus ultrn). 304 (green, the n Violet wolling b 0) 0 304 (violet kolunie (inon plus ultrn). 305 (violet kolunie (inon plus ultrn). 306 (violet kolunie) 306 (violet kol	Æthyl violet 305. Green, bluo, then violet Benzyl violet 6 B, 7 B 1305. Green, bluo, then Raris violet 6 B, 7 B Methyl violet 6 B, 7 B 1306. Kreen bluo, then Violet B B, 7 B Violet B B, 6 B, 7 B 1306. Violet B, 100 C	Acid violet 4 il N. 6 B. { 312. { then g reen, held violet 7 B, N. Acid violet 6 B. 316. { line-green. hine-green. Formyl violet 8 4 B. { First olive-green. then green, then green hine. Acid violet 4 B extra { 817. } hine. green, then green plt. Alkali violet 4 dirty green plt.

VIOLIST - Consinued.

Ppt. Ppt. Ppt. Ppt. Ppt. Ppt. Ppt. Ppt.	Red-vlotet ppt. Ppt. Ppt.	Hino, then green, with excess deep yellow-brown,	364. Then yellow,	With excess deep	agr. Brown, on dilution		(Bline sof, and do, ppt., then green, then brown-vellow sof.	i Ppt, with excess, yel-
Vroteer Fontlaurel, Oxamino violet. Azamanyo II. Azamanyo II. Congo cacinti III. Congo cacinti II. Congo cacinti III. Con	Hear Folds reddelt 148. A zo vlotet 1900 (200) Heading vlotet 2000	Methyt violet 14 and 2 11. Methyt violet V 3. Violet de Ducla. Violet an methyt 14. Violet sednide (non plus tillra).	Organial wholet a trop of the Chystal wholet B B O. O. Violet C.	Benzyl vfolet Brand Brand Market Mark		Violet Impérint (phenylique)) Aela violet 4 B N, 6 B	Povinyl violet, 8 4 B	Alkall vlolet, 314, i Ppt. with exceen yel-
14th Mue-red s o l., then Alls. Furbalue red, 4Hb. Furbalue red,	462, Pine, Heat violet red. Harm Page 1984 19			Solution	Violet-black ppf.	Pres color nold ppt.	Ppt, in cene, sot, Ppt, Blue-violet ppt,	Ppt.
4 H. 4 H.	Mentiforne virter il RA Fuchsin, Resulent	No Unactron; Red Violet fi B extra. Violet H N, 5 B, B, B E. Gallefin in paste.	Allarine violet	VIOLET. On Addition of IGB to Water Solution	Dradk; Diamond vlulet N.,,180, Vlotet-black ppl.	WHITE: Obrame violet	v (ofet-blubet) 183, o earlinth (1) 183, drope (2 B) 185,	Jordenius extra Comgo violet. 1845. Ppt.

HLUE-Continued.		YELLOW:
Fast acid violet B. Violamine B.	Flocks.	Hofmann's violet
Fast neutral violet II	With ехсень.	5 В О, О
Methylene violet It It A 462.		Athyl vlolet
Clematine. Sufranline M N.		Fast acut violet 1015
Heliotrope B. 2 B	With large excess.	Wool violet S
Amethyst violet465,		Azo acid blue B
Rosolan 466, On large excess first	On large excess first	Alkuli violet
Paraphenylene violet	Ppt.	Acid violet 6 B N
GREEN:		Fast acid violet A 2 R
Hofmann's violet302.		Gailo cyanine D.H., B.S
Acit violet 6 H. 316. Bine-green. 120. Grass green. Rosdan. 466, {On large excess, first	Blue-green. Gruss green. On large excess, first	Princ pure.
BROWN: (Ding, the	Conne, then green. Red-bround	No beachton: (Chromoteul B
Red violet 5 R extrn		Red violet 6 R S
Violet 4 R N, 5 R \ 302, Yellow-brown. Violet R, R R.	Yellow-brown.	Red violet 4 B S
Regina violet.	On dilution blue.	Neutral violet
violet imperial (puenyhque)) Acid violet 6 B N	Violet, with excess	Heliotrope B, 2 B. Tunnih heliotrope Cacess Trumh heliotrope Care Care Care Care Care Care Care Car
Gallein in paste	Yellow-brown,	Rosolun 466, With large excess growing first blue, then green.

VIOLET-Continued.

00		1 12.	AIL	-ъ	MOK	r O.	1, (COL	OKI	.O.L	S AI	ND D.	LEI	ıo.	
	Gray-violet ppt., on warming colorless.	Blue-violet ppt., on warming colorless.	Red-violet.	Ppt, of base,				467. Blue-violet ppt.		•		Becomes bluer and brownish ppt.	Flocks, on warming colorless.	Dil., light blue; conc.,	Flocks, on warming colorless. Ppt.
VIOLET—Continued.	Althyl violet	Acid violet 7B	Gallocyanine D H, B S	Rosolan 466. Ppt. of base.	Mauveïne Rosolan Violet paste	Chrome violet	AnileineAniline violet		Indisine Phenamine	Phenameine	Tyraline. Tyrian purple Lydine	BLUE: Fast violet reddish148. Becomes bluer and	Acid violet 4 B N	Acid violet 6 B316, Dil., light blue; conc.,	Formyi violet S 4 B
					467.					ater Solution.	96. Blackish ppt., red-	36. Blue-violet ppt. 51. Ppt. in conc. sol. 88. Ppt.	12. Dirty violet.	49, Blue-violet. 96. Red-violet, blackish	04. Ppt.
No REACTION—Continued.	Manveïne	Violet paste,	Manye dye	Antline violet		Indisine Phenamine	Purpurine. Tyraline	Tyrian purple	VIOLET.	On Addition of NaOH to Water Solution.	Brack: Chrome violet paste	Violet S. 36, Wool violet S. 36, Ract violet bluish 151, Camilia Volet. 188.	Azomanye B	Hessian violet	Crystal violet 5BO. O 3804. Ppt. Violet C

Thrns redder, Cherry-red, Fuchshie red, Brown, red ppt,	303, Brown-red sol. and ppt.	306. Srown-red sol. and ppt.	462. Brown-red ppt.	Dark red ppt., on di- lution soluble with red-violet color.	464. Ppt, soluble in water.	On excess partial ppt.
HED — Continued. Heliotropie 2 B. 185. Congo corluth H. 214. Azo vlolet. 233. Hofmann's vlolet 302.	Methyl violet B, 2 B. Methyl violet B, 2 B. Violet do Paris. Violet an methyl B. Violet an inthlyl B. Violet soluble	Benzyl vlolet T B. Artis vlolet B B. T B. Methyl vlolet B B. T B. Vlolet E B. T B. Vlolet E B. F B. Vlolet au méthyl 6 B.	Methylene violet R R A Methylene violet 3 R A Fuchish	462.	Heliotrope B, 2 B. Tannin heliotrope.	No legaction: Chromotrope 8 B. 100. Chromotrope 8 B. 100. Diminie worder N. 180. Heliefrape B. 232. Fast acid violet B. 314. Fast acid violet B. 348. Fast acid violet A 2 R. Volamine B. 349. Fast mentral violet B. 349. Fast mentral violet B. 349. Violet mentre solide B. 349. Amethyst violet . 465.
	Ppt., sol. turns bluer, Fpt.	que)	Ppt.	- br.	Gray-violet, ppt., on warming colorlens, leaving a brown	Mire tlocks, on warming colorless. Dit, light blue; cone., colorless. Blue flocks, on warming clorless. Slowly bleached, lillne-vlolet put., on warming colorless. Yellow-ved.
BLUE—Continued. Gallein in paste. Gallein Am paste. Alizarine violet. Anthrucene violet.	Huows : Fact violet reddish 148. Ppt., sol. turns bluer, Red violet 5 k extra 148. 148. 149. 1	Regina violet Pat. Pat.		COLORERS	Æthyl violet,	Acid vlotet 4 B N, 6 B, 7 B N 312 } Interflocks, on warm-Acid vlotet 6 B 316 } Interflocks, on warm-Brand vlotet 6 B Formyl vlotet 8 4 B \$17 } Interflocks, on warm-Acid vlotet 6 B N \$17 } Interflocks, on warm-Acid vlotet 6 B N Acid vlotet 7 B \$19 Slowly bleached Acid vlotet 7 B \$19 Slowly bleached Acid vlotet 7 B \$10 Slowly bleached Acid vlotet 7 B \$10 Slowly bleached Chromotrope 10 B 63 Yellow-red Congo corinch 6 \$3 Yellow-red Congo corinch 6 \$183. Cherry-red

Matte = Continued. Matte = Continued. 238 Green-blue. Matter stars to the continue stars to the continue stars. Matter stars to the continue sta	Naphthyt blue 2 H. Tenzo bluek thin R. Tenzo bluek thin R. Tenzo hidge blue 201.	Alkarine eyanine R in paste,	>	Authracone blue W B	Coleste blue B	Ultertocyanine 422, Ulter slowly, Oldlinghe A. 421, Owering A. 422, Owering A.	Fluorescent blue	First bling R. & Radondos saturble) Solid bling R. & Radondos saturble Industries advanced and advanced as a resolution of a contract	Azhronae reman sonton. Underno P. F. Undephonic extra.	Printing Pine H	Printing blue it & B	Nofr a Patenol Complere blue	Note O N N
Blue gray.	Red-vlotet,	Red vloter.	Red-vlotet,	Brown-ylolet,	-158, Green-blue.			Oreentah blue,	Clear blue, Dark blue		Bluck-blue,		
DACK: Metaphenylene blue B		Gallanti findigo P	197.	Neutral bluedbb. Brown-y-blief	mine blue B B.	Azo bluck blue. 211, Azo blue	Diamine blue II X. Henze blue II X. Congo blue II X.	218,	22.25. 22.25. 22.25.		Noir blen Diamine E.	Bengal blue G. Catten blue G.	Benzo azurine 3 (1

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Contin	
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UE	į,
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	430. Dirty green.	434. Blue-green.	Yellow-green.	440. Yellow-green.	Yollow-groon	Yellow-green.	ынскв ъг кен, Вис-ктесп, Ипс-green,	Greenish-brown,	Red-brown.	Yellowish-brown.	Red-brown,
GREEN-Continued.	New bine G. New bine G. New bine G. Net bine 2 B for cutton, Metamine bine B. G. Fax catton bine B. Pax may bine B. Naphthol bine B.		Gentlanine	Methylene blue B II, extra D. Methylene blue B II, in powder extra. Methylene blue B II, in powder extra. Methylene blue B B extra. Methylene blue B g erystals. Æhlylene blue	Thiouin blue G. O extra442. Thioenraine R. R en pate443. Toloidine blue O	New methylene blue N	Nuphthazine blue	Brown: Tadofine blue R 73.	Methyl alkali bita. Alkali bita 9 Alkali bita 61 Alkali bita 61 Alkali bita 61	Bavarlan blue D S F	Methyl blue for cotton Methyl blue for cotton Brilliant cotton blue greenish Methyl water blue Cotton blue
						Blue-green.	Black-green, Black-creen,			Blackish-green.	
BLUE—Continued. Frat blue 18	First blue 3 R. Hen solide R. 3 R. Fast blue 4 B. Fast blue 6 B. Fast blue 6 B. Solid blue R. & B water soluble.	Solid blue 4.5. Blue C. & B water soluble	Induline B & B.	Nigronaeus water solluble. Gray R. & B. Blett de Bengale. Paruphenylene blue R. Fast new blue for cotton.	Toluylene blue B for cotton	Diamine pure blue.	Congo jure blue	Capri blue G N	New cotton solid blue. Naphthylene blue R in crystals Fast blue R for cotton in crystals	Fact may blue Fast cotton blue R, R R, 3 R crystals. >429. Bluckish-green. Blen pouroun P.	Fast blue III R. Fast nay blue R.M. M. M. Naphthol blue R. D. Frat line for cetton. Meldolas blue

BLUE - Continued.

2001. Brown-yellow,	315, Brown yellow.		. Bu, Durk ved şellaw	. 3394.	#35. Brown-yellow,	illi, Bark yellow.
Vectors Continued. Clearlin blue ff ff. Abedian blue ff ff. Oped blue F C Oped thine R F C Hessian blue ff. High blue Light blue High cholos southle	Fast wood blue. Alpha blue. \$115, Heaven yellow.	Water blue 41 Bester. Water blue Chim blue Chim blue 8 H. Heut de Lyeur. SHE blue.		Victoria blue B A alcohol soluble	New Victoria bine	Authricene blue B Albachie blue Soluble powder A II B.
-224, Bed-brown,	Red brown,	Yellow brown, Yellow brown, Red-brown, Yellow brown, Turoling blue later on Red-brown,	Uresulab-brown, Byolutlan of lodine on warming,		Vollowish, Brown yellow,	Hrown yellow.
Hnows — Confirmed. Boverion thus D. B. F. Helvedle to the Residual to the Southle flow B. H. Southle flow B. H. Southle flow B. H. Diptropylamine blue. XI. Southle thus Hiet methyl. Pure blue B. B.	Alkall blue N G. Soluble blue X G. Soluble blue X G.	Night thine Visitation thine Authorsees blue W (1, 140) Ladeschool in provides Flucture yadding the C (1, 140) Flucture yadding the C (1, 140) Illusture yadding the C (1, 140) Illusture halp plane in one C (1, 140) Illusture the C (1, 140) Illust	Goldentine H. Control of the C	Acid blue 6 G.	Fatesti bine V, N, Superflue, conv., (202, Cymple of N, O, R, 202, Cymple of N, O, R, 202, Fatest bine A, 201, Cymple bine A, 201, Cymple of Cympl	of solutible Calcoliol solu alcoliol soli

Ppt. 1'pt.	Fig. Blue-violet ppt, Ppt, Brok. Block-violet ppt, Prock-violet ppt, Prock-violet ppt, Prock-violet ppt,	ed-violet sol., then blue flocks.	rty violet, then blue,	pt ater vlolet, then volet plut. irst blue, then blue- violet.	First green, then violet ppt.
	236. 236. 237. 237. 257. 267.	Fast neid blue it Violamine 2 it Fluorescent blue 123. Reid-violet sol, then Fluorescent blue 123. Later red-brown ppt.	New blue B. Fast blue 2 B for cotton. Fast blue 2 B for cotton. Metamine blue B. G. Fast cotton blue B. Fast cotton blue B. Fast nay blue H. M. G. M.	.433. F 434. { .439. F .446. E	
YELLOW—Continued. Nile blue A	Alkali blue. Nicholson blue. Fast blue Blue Medilin. Hoechet new blue. Cirone blue. Sigh. Brown-red. Sigh. Brown-red. Sigh. Brown-red. Fast said blue blue. Sigh. Brown-red.	Violamine 2 R	Alizarino bine A B	Violett: Violett: Indoine blue R. Dlamine blue B. Benzo blue B. Congo blue B. 138 189.	Azo black-blac. 211. Blue-violet ppt. Azo blue. 216. Ppt. Diamine blue II X. 217. Ppt. Congo blue II X. 217. Ppt. Diamine blue II X. 217. Ppt. Congo blue II X. 2218. Ppt.

BLUE-Centinued.

Ppt.	1		Later py t
	, , , , , , , , , , , , , , , , , , ,	÷.	325. I
	Methyl blue for silk M L B. Navy blue B Methyl blue Methyl blue for celton. Methyl blue for celton. Mellinat celton blue greenlett. Mellinat celton blue greenlett.	Action blue Methyl cotton blue Methyl cotton blue If wordin blue D B F If wordin blue D B F If wordin blue B B. Solnible blue B B. Diptenylamine blue. X L wolnble blue	Fure bine B S I Water bine B S I Water bine 6 B extra Water bine 6 China bine China bine China bine Silk bine Bine astra Cotton bine B S Water bine B S Water bine bine Loadon bine extra Cotton bine B Water bine bine Loadon bine extra Belu marine
		ish-blue at. et. ish-blue	ı, then
	Ppt. Ppt.	237. Clear blue. 238. J Dark reddsh-blue 245. Violet blue ppt. 247. Dark greeniel-blue	2011, J. P. Prict. green 308. Pric. 309. Pric.
176.		28.5. 28.5. 26.7.	
Volley—Cuntivued, Bold bine R R and B water soluble. Sold bine B water soluble. Hue C B water soluble. Induline R and B. Induline 3 H. Nigroshio water soluble. Gray R and B. Bleu de Hengale.	BLUE : 227. Diamine blue-black E . 228. Noir blen	Diamine pure blue 217, Cleur blue. Part, Cleur blue. Part, Cleur blue. Part Part	Cyanol extra Acid bine Green, then Acid bine Green, then Bavarian bine alcohol soluble. 300. Prt. Light bine superfline alcohol soluble. 300. Prt. Light bine superfline alcohol soluble. 300. Prt. Alcohol bine Green, then the first superfline alcohol bine 400 bi

Later blue-violet,	Clear blue, Dark blue,		Light blue ppt.	Later violet ppt. Ppt. First green, then blue.
BEC		Fast blue R and Halcohol woluble. Sold blue R R and Halcohol soluble. Induline alcohol soluble. Azine blue alcohol soluble. Indigene D. F. Indigene D. F. Indigene O. Renhol soluble. Prinching blue H. Prinching blue H. Prinching blue H. Vireche alcohol soluble. Voir A' fulcool Complete blue. Complete blue Nigreche alcohol soluble. Nigreche alcohol soluble. Nigreche alcohol soluble. Noir A' fulcool Complete blue Noir A' fulcool Complete blue Noir C N N.	Para blue478. Light blue ppt.	GREN: T3 Later violet pat. Intofine blue R
DECENTATION PRINCE PRINC	Frat soid blue R. Sept. First red violet soid	Fast navy bine. Fast cotton bine II, IR, 3 R crystate otton bine II, IR, 3 R crystate bine. Thenylene bine. Hen nonvean P. Fast bine III II. Fast navy bine IB. M. H. M. Naphilol bine B. M. H. M. Naphilol bine B. M. Naphilol bine B. M. Naphilol bine B. M. New bine G. Fast bine G. Fast bine B. Fast bine bine B. Fast cotton bine B. Fast cotton bine B. Fast navy bine it M. G. M.	Naphthol blue B	New Michylene bline G G (44), FFrat green, then Greens. Nile bline A delay bline A delay bline. Muscarine Advantage Campanuline. (434, Later vlote, then Cyan Campanuline.

BLUE—Continued.

	Yellow red.	Fuchshe red.	Yellow-red.		Solution.	Ppt.	Blue-violet ppt. Ppt.	Ppt.	Bedder, on excess Violet ppt. Ppt.
→ 427.	3865	45. 416. 12.	484.	. } 476.	Vater	189.	, 211. , 216.	₹217.	} 218. 219.
$\begin{array}{llllllllllllllllllllllllllllllllllll$	Altarine blue in paste	Aeld alizarine blue II B and G R412. Capri blue G N	Gallanii indigo P. S. 424, Yellow-red. Coreino. 426.	No Reaction: Paraphenylene blue It	BLUE. On Addition of HCl to Water Solution.	VIOLET: Diamine blue B B	Azo black-blue. 211.	Definition of the D. X. Congo blue B. X. Congo blue B. X.	Johnnie blue 3 B. Reaco blue 3 B. Congo blue 3 B. Oxamine blue 3 R. 218. Redder, on excess congo blue 3 B.
lfonic	ехеевя , then		., then						green,
Later ppt, of su acid,	First yellow, water green, blue, Light green,	Later blue. Later blue. Later blue.	Later blue. Later violet sol	Red-brown.	Ppt.	Ppt.		Later green.	Later green. Excess water then blue.
.291.	331. 331. 335.	336.	432.	.339	393.	-4 1 4	} 452·	292.	.293. { 334. {
GREEN—Continued. Patent blue A	Apaic Wool blue Apaic Mool blue Victoria blue B 331 Stret yellow, excess Victoria blue B 331 Stret yellow, excess Victoria blue B 331 Stret yellow, excess Victoria blue B 332 Stret yellow, excess Victoria blue B 333 Stret yellow B Stret	Night bine Night bine Victoria bine 4 R. 337. Later bine. New methylene bine 6 G.	Nile blue A	Baown: Chrome blue	Alizarine blue S in powder, also in paste. Authracene blue S	Alizarine blue Soluble powder A B S	Colorless: Cynnine Chinoline blue.	YELLOW: Patent blue V, N, superflue, conc 292. Later green. Reton blue 4 B N, G, R	Cyanine B

	Ppt. Ppt.	Ppt Ppt Ppt		315. { Blue ppt. and green sol. Ppt.	Ppt.	Darker blue.		1994. Little HCl no reac-	
53	227.	8 8 8	264		355	328.		#	
BLUE: Indoine blue R	Diamine blue B. Diamine blue-black E. Noir bleu.	Diamine F	Benzo black blue 5 G955. Benzo black blue 5 G267.	Pact wool blue Alphe blue Methy alkali blue Alkali blue D. Alkali blue b.	Alkali bite Nichisen bite Fast bite Illeu alcalin	Bavarian biac D S F. Methy I biac water soluble. Methyl biac for silk M f. B. Navy biac B.	Methyl blue. Methyl blue for cotton Brilliant cotton blue-greenish. Methyl water blue. Cotton blue. Methyl setten blue.		Pure blue is S 1
Ppt.	Ppt.	Blue-violet ppt. Illue-violet ppt. Ppt. Blue-violet ppt.	Gallanii indigo P. 424. Red-vloter. Gallanii indigo P. S. New blue R.		429. Later blue.		Dirty violet,	Nile blue 2 B	Toluylene blue B for cotton
233.	235.	Naphthyl blue 2 R. 256. Naphthyl blue 2 R. 257. Brazo black-blue R. 261. Benzo indico-blue R. 265.			ह्यं		-130.	Nile blue 2 432, Nile blue 2 433, Nile blue 2 1, 433, Nile blue 2 1,	-1
	Benzo azurine G. Bengal blue G. Cotton blue G.				<u> </u>		New Oute to Netramine blue B G. Fint cotton blue B. Fint cotton blue B. Fast may blue B. M. G. M.	Nile blue A 432, 1836 1842 183, 184, 183, 184,	7

BLUE-Continued.

Alcoholie salution Al- i most bine.	475. Deeper blue,	Light blue ppt.	Dark green ppt, l First green, then yel- l low. Green, then yellow.	Green, then yellow. Ppt. of sulfonte acid. Hine ppt., green sol. First ppt., then green, then durk yellow- brown sol. JYellow-brown, on di- htton green,
Brus—Continued. Noir à Paleool Gonțiera blue Bleu Compiera l'Inlevol Noir C N N.	Frast blue R. Faat blue S. Ban solide H. S. Blon solide H. S. Brast blue H. Frast blue B. Frast blue greenist Folid blue R. R and H water soluble. Solid blue R. R and H. Indulline G. B. water soluble. Indulline R. B. M. Chippenen water soluble. O'Igreene water soluble.	Men de Bengale Para blue	Ornera: Firm bino. Symmetry. Cymnel extra. Acid bine if 0., superfine, conc	Oxonine II. Oxonine II. Fateur blue A. Alphe blue. Victoria blue II. Victoria blue II. New Victoria blue II. New Victoria blue II. New Victoria blue. Victoria blue II.
127. Ppl. 51. Flocks.	420, i First violet, then		faft. Ppt. 18th: Slightly bluer. 7tl. Deeper blue ppt. 17tl. Ppt.	471. J Mechalle solution al- inost blue,
Brune—Contenued. Alkall blue X G. Soluble blue X G. Frat acel blue R. Volomine 2 R. Froton	New blue R. Naphthylene blue R in crystals. Naphthylene blue R in crystals. Fact blue R for catton in crystals. Fact blue R for catton in crystals. Fact nary blue. Fact catton blue R, R, S, R, crystals. Flooring blue R, R, M,	Meldolar blue O	Basel blue B B In parte. 450. Meta phenylene blue II. 460. Naphthazine blue 1. 471. Foiling blue 471. Fast blue R and B nicohol sentuble.	

	M. and and loss			. Pht.				. 3ft2, Yellow-red.		Turns slightly red-		Fuchshe red.	. Ppt. and red sol.				In alcoholic solution.
YELLOW-Continued.	Alizarine blue S in powder, also in parties and partie	Allzuriech ing S. Soluble powder A B S	Corontess	Coreino A B.	Cynuln	Chinolin blue	RED: Allzarine blue in pasto	Alizarine blue G W 8992 Alizarine blue A N F, R, B R Alizarine blue A N J F, R, B R	Capri bine G N	Delphin blue	Gallamin blue430.	Celeste blue B	Gallozino A	No heaction: 3 18 936	Diamino puro blue.		Gentlan blue 6 B. Alcohol blue. Alcohol blue S. F. C. Opul blue S. F. C. Bluel bluider Heel bluider
_	First ppt., then green, then yellow-	First ppt., then	brown sol.	First ppt., then	yellow-brown sol.	Yellow-brown, on di-	_	First ppt., then green, then yellow- brown.	Red-brown,	To alreholic sol., red-	Yellow-brown ppt.	Оп ехеекв.		First green, then yellow.	Green, then yellow.	Green, then yellow.	Red-yellow.
GREEN - Continued.	Night blue	Victoria hino d B		orla blue B	Victoria bluo B S alcohol solutile ("")	New Victoria blue	Night blue	Victoria blue 4 B	Chrome blue	Indephenel in powder [114.] To alcoholic sol., red-	Finorescent blue	New methylene blue G G	YELLOW:	Cyaniol extra	Patent blue V N, superflue, conc 292. Green, then yellow. Keton blue 4 B N, G R.	Cymlue B	Alizardne blue lu paste. Alizardne blue ft. Alizardne blue G W. Alizardne blue A. D N W. Alizardne blue Ft. R. R. R. Alizardne blue Ft. R. R.

BLUE-Continued.

No object. (124. On excess time put. (125. Partial blue put. Vro		m large excess, slightly bluer.	vater solution. 73. Blackish-violet ppt. 189. Red violet. 218. Herk violet. 219. And violet. 220. Bed violet. 220. Red violet. 230. Red violet. 230. Red violet.	On warming. On warming.
Exactions—Continued. Split blue activities or solution. Split blue activities or solution. Split blue activities or solution Split blue activities Ac	No Heactron—Continued, denibanh Methylene blue H H, extra D, Methylene blue H H, extra D, Methylene blue H H in powder extra, Methylene blue H H in powder extra, Methylene blue H H extra. Methylene blue H I explais.	Titionin blue Q, O extra 492. Titionin blue R. 1948. Titioarmine R. 1948. Titioarmine R. 1948. New methylene blue N. 465. Induzine M. 1948. Puraphenylene blue R. 465. Puraphenylene blue R. 476.	On addition of NaOH to water ine blue It 73. The blue II 73. The blue II 18. To blue II II 8. To blue	204.
The Accritist — Continued, Rich laine The alcould soluble Totally laine Totally laine for solun Totally laine for solun Totally water blue Totally blue Total blue Totally blue Total T	In alcoholto solution.	Он ехееяв Ине ррб.	Partial blue pyt.	Он ехсеяя Іноми,
THE AREAUATIES AND PRODUCE OF THE AREA	No Reacross—Continued. Light blue Har atomid soluble Blen conde. Methyl blue Methyl blue for cotton. Refillmet cofton blue greenlelt Methyl water blue.		Water blue ff B extra Water blue ff B extra Water blue for the first blue for for first blue from the first blue ff Water blue B for for for first blue for for first blue for for first blue for first blue first blue for first blue first blue for first blue for first blue for first blue for first blue fir	Acid alzarine B it out O R

		In alcoholic sol, dirty red or red -violet sol, and dito ppt.			455. Brown-vlolet nut.			Ppt.	Brown-vloiet ppt.	Reddish-blue. After a time blue ppt.	Оп ехсевы дтееп.
Violet + Continued.	and B	Ĕ	First blue R.	Fact bine 8 ft. Blue solide ff, 3 ft. Fact bine 8. First bine 6 ft. First bine 7 ft.			Induline 3 B. Nigrosene water soluble. Gruy It and B. Bleu de Bengalo .	Paraphenylene blue R	Para blue	BLUE: Diamine bine B	Alizardine bitte in paste. Alizardine bitte it. Alizardine bitte it. Alizardine bitte A, D N W F, R R II. Alizardine bitte A B. Alizardine bitte A B. Alizardine bitte A B. Alizardine bitte bitte A B. Alizardine bitte bitte A B. Alizardine bitte bitte A B.
On warming		Blue vlolet.	Ppt, (Red-violet, on dlu-	tion red flocculent ppt. Blue-vloiet, redder on the edges.	Blue-vlolet.	Turns deeper violet first, then dirty violet ppt.	Deeper vlolet, on ex- cess firty vlolet p, t.	On excess put.		Dirty-violet ppt. Ppt. Ppt.	In alcoholic sol. dirty red or red - violet sol. and dito ppt.
VIOLET - Continued, Patent blue A	Fast acid blue R. Violamine 2 R.	Alizarine thue 8 in powder	Capri blue G N	Unliamme bino	Gailanli indigo P. Gailanii Indigo P. S. S. Blue-violet.	Gentlunin 439.	Methylene blue 04, 81 (1, 1, 1, 1, 1, 1, 1, 1, 1, 1, 1, 1, 1, 1	Athylene blue G. O extra.	Thiocarmine R. Thucarmine R en pâte.	Toluldene blue O	First Direc R and B alcohol soluble Solid bine R L& B alcohol soluble Ludulitar alcohol soluble Azhre bluke alcohol soluble Indigene D, F. Indipperime extra Blue C B alcohol soluble

. 32 1. Red-brown,	Dark red-brown ppt.	Flocky ppt,	Pale red-brown ppl. Violet-brown ppl. Turning bine on		Ppt.	
Methyl blin Methyl blin Methyl blin for colton Nettyl blin for colton Irilliant estron blin greenish. Methyl water bline Certon bline Certon bline Isvarian blin D IF Heveria bline 8 II Soluble bline 8 II Soluble bline bline I liphenydamine bline X is coluble bline III methyl bline III liphenydamine bline	Pure blue is 8 l	New Victoria blue L., Fl35, Flocky ppt,	Night titte	Now blue R. New extron solid blue Naphlityfene blue R. in crystala. That blue R. for cotton in crystala. Colton blue R.	Fast may time Fast outon bine B, B, B, B erystelle, Phenyleine time Hent neuvon P	Fast invy blue B. M. M. M. Naplikiol blue R. D. Fast blue R. D. Fast blue Fast blue Fast blue Mediciae blue.
	Hue-bronze ppt., turns brown on warming.		ли2, си ехсеня,	221, Red-brown.	1122, Hed-brown.	Hed brown (ef, red regetlen).
Alzarine eyanine R in paste	Oyunlue	Organic blue in paste		Mony s. Methy blue. Methy alkell blue. Alkell blue fr. Alkell blue fr.		Mayarlan blue 19 S F. Methyl blue water soluble Methyl blue for silk M L II. Navy blue II.

RED—Confinued. Benzo azulue G Benzo azulue G Benzo azulue G Benzo blue G Cotton bl	Cyanol extra. Cyanol extra. Acid blue 6 G Gentian blue 6 B Alcohol blue 7. Alcohol blue 8 F Alcohol blue 8 F	Opal blue. Hen hmière 309, {In a le o li o 1 l c sol. Hestin blue. Light blue. Blue alcohol soluble. Bleu quide.	Bavarlan Bito D S F. Methyl bine water soluble. Methyl bine for silk M L B. Navy blue B.	Water bine 6 B extra. Water bine 6 B extra. China bine. Cofton bine 3 B. Blea de Lyons. Silk bine. Hare extra. Cofton bine B S. Water bine B S. Soluble bine. London bine extra. Blea marine.
٠,-	New methylene blue N	Heack: (Vlote - black sol. & diop pic, on warm-Alkall blue X G	Now methylene blue G G	Coloutress Col

BLUE-Continued.

	On holling violet. On boiling violet. On boiling violet.		that; coor returns On facility differ. Silghily differ.	
No Beadwitton : Diambine blue-bluck E Solr blen Standine E Henzo bluek-blue & G 887.	Facint bline V, N, auparine, cond. (2012) Keton bline G, R. Cyanline W. Facint bline M. Facint bline M.	Post wool blue Alpine blue Offenne blue fording the powder fording the	Irla blue Thiomembro R. R en pfte. 443. Napithazine blue. 443.	
96.	tion red document. 1914. [NaCH (1 : 1000) 195.; 4 th or extraction given brown ormage and with dark circum	dimeacence. Brown-red ppt., sol, in other with yel- low-red nelor.	Alcohollo sol, diriy- red or red violet, ad and discont.	
avb. 325.	4:12.		474	
REGIN - Continued. Pure blue. Fure blue. Hown red.	Chillantina blue,	Nilo bine 2 H. 43H. Facellot moltible	A Attin blue attorned sections of Atting blue attorned sections of a Independent of the Independent of the Independent of the Attorney of the Independent of the Attorney of the Independent of the Attorney of the Independent of Independent	Fritting thin it and B. Acedine blue. Nigressure alcohol soluble. Noir A Talesol Couplers thin. Hen Coupler's Talesol Noir O N N.

GREEN. On H₂SO₄.

BLACK:	6. Ppt.; evaporation of 112S.	Yellow: Malachite green cryst, or powder Malachite green B	<i></i>
Brown: Azo green Cerrulcine in puste	7. Red-brown. 7. Dirty yellow-brown.	Benzal green New Victoria green New green Solid green crystals	
Carnleine S in powder or in paste 368.	8. Dark brown.	Solid green O. Vert solide eristnux O.	
Gumbline R in paste376.	6. Red-brown.	Dismond green B.	
Gambline († in paste) Gambline X in paste Gambline Abace green J	377. Dark brown.	Fast green Brillant green Malachite green G.	
Naphthol green B). Yellow-brown, 8. Brownish.	Athyl green. Smarrigd green. Solid ages I	285.
VIOLET:		Solid green J J O	
Delizo olive 262. Unannine green B 268. Vool green S 338.	s. 8. Brown-violet.	Victoria green 3 B	286.
Fast green M435.		Guinea green B288.	.288
Alizarine green B	7. Dirty blue-violet. 9. Blue-violet.	Light green S F bluish	080
Blue:	(Rine or red sol and	Acid green M	
Acid alizarine green B & G413. (strong fluorescence.	s. t strong thorescence.	Light green S P yellowish	
Green: Diamond green	3. Blue-green.	Acid green D	290.
Dioxline.	*	Vert acid J J extra concentre	
Alizarme green G438,	i. Illue-green.	Methyl green	
Methylene green extra yellowish G l 441. Dark green.	l. Dark green.	Double green S F. Light green.	7328. Rea-ye

GREEN-Continued.

XELLOW—Continued. Akthyl green. Mathyl green. Ona.nor. Chrome green. Chrome green. 205. Yellow-orange.	Yellow-orange,	Bhown—Continued. Gambline G in pucke. Gambline X in pucke. Gambline. Alance grieci J.	K8,
Find green extra 2918. Park green extra 2918. Park green extra binini.	Yellow-red.	Azing Green Green Azing Fleet Incovability then Azing Green Az	t brownish, then ne-green.
391.	With development of gae, cherry-red. Bine or red sol. and strong fluores-	Violer: Diamond green	First bine-green sol., excess water bluck- violet ppt,
Gallauli green (1	Carmolaine red.	Alizarine green G436, { Pira pi	First bine, then vio- let sol., then red ppt.
GREEN. On addition of H ₂ O to H ₂ SO ₄ Solution.	solution.	BLUE: Pirate Pira	First green-bine sol., excess water black-violet upt.
Denzo olive	Green-black ppt. Black ppt. First reddlal-brown, then greenish sel., & green-black ppt.	Fast green extra binlsh 298, confined green extra binlsh 298, confined green G 298, confined green G 298, confined green G 298, confined green G 298, confined green extra yellowlsh G 411.	Almost colorloss, ex- coss water green- bino. First bine, then vio- ict sol., then red ppt.
vwn: Cœruleine In paste	Reddish-brown, then greenful sol., and green-black ppt. Yellow sol. and brown ppt.	Greek : Diamond green	First green-blue sol., excess wuter bluck- violet ppt.

.367. First reddish-brown, .367. then greenish sol.,	.391. Stree chinoline, resp. sulphate.	436. First blue, then violet	437. First yellow-green sol., then red ppt.	458. First brownlsh, then 559. blue-green.	(Almost colorless, ex-	298.) cess water green-		-	excess water green.	(First red.yellow, then	green, men
(IREEN—Continued. Cornleine in paste	Altzarine green S391.	Alizarine green G	Alizarino green B437.	Azine green G B458. Azine green S459. Integreen S459.	Colorings:	Fast green extra blulsh	Yellow: Malachite green cryst, or powder	Malachite green B. Benzal green New Victoria green New green	Solid green oryst. Solid green oryst. Vert solide eristaux o. Unmond green B. Find green B.	Brilliant green G. Malachte green G. Dianond green G.	Arush Breen Smirragd green Solid green J J O
	Pirst dark-yellow, then yellow-green.	(excess water green,		(First red-vellow.	~	286. Red-yellow, e x c c s s	First yellow-red, then yellow-green, then	- 33	290. Slowly green.	328. Yellow-green.	329. (Yellow, excess water green.
JBEEN—Continued. Malachile green cryst, or powder Malachile green B. Henzal green		Solid green O	Diamond green B.	Brilllant green. Malachite green G.	Athyl green. Smaragd green J.	Victoria green 3 B	Gulnen green B258	Light green S F bluish Acid green M Acid green M Acid green M		Methyl green Vort čilncelle Double green S F. Light green	Æthyl green 1 329, { Yellow, excess water Methyl green 5 329, { Freen. Wool green S 338. Yellowish-green.

GREEN. - Continued.

GREEN. On Addition of HCl to Water Solution.	HLACK; Dlamine green H	Italian green	Achi green A bhish Achi green M	Acut green minet. Light green 8 F yellowish. Acid green extra conf. Acid green extra conf.		hrown.	Violiste: 168. Divine reddibilisheletet	Brane:	Acid alizaring green II and G	Solid green G 420, Dark bine ppc.	Chicken;	Victoria green 3 H	New world green 1: ft mid 3 11	Double green S F (100 yellow-green)
YELLOW - Continued. Victoria greet H H		Light, green S. P. blutshi. Aeld green	Acht green minen Althyl green Nathyl green Achtyl green Achtyl green Achtyl green Achtyl green	133.	Naphthol green Π 339 , g_{13} , and K_{4} Pec $(y_{21}, y_{122}, y_{133}, y_{133}, y_{133}, y_{133}, y_{133}, y_{1333}, y_{1333}, y_{1333}, y_{1333}, y_{13333}, y_{13333}, y_{133333}, y_{133333}, y_{133333}, y_{133333}, y_{1333333}, y_{1333333333}, y_{1333333333333333333333333333333333333$	ONANGE : 435.		Quinea green II	~	Oloxhue	13.	Alizarine green G	Allzarine green H,	No REAUTION: Olivone green

OHANGE: Chrome green	Azo green	treen B437.	No Reaction: Ceruleine in paste	Gambine R in paste	Abace green J	GREEN.	Un Audición of Anul to water Solución. Brown: Reizo olivo	VIOLET: Isku green SF bluish		Acid green Acid green 290. Colorices and turbid Acid green 200. Violet ppt. Verta coloritie 200. Verta coloritie Verta colo
	Turns greener. Ppt.		Red-yellow.		Red.yellow.		Yellow green, then yellow. Brown-yellow.	Yellow, excess water greenish-yellow.	338, [Red-yellow, on dilu-	First green, then yel- low. Brown-vellow.
320.	458.	_	 - 381.		- E55.		286.	2888	358	1 Sign
GREEN - Continued. - Rehyl green. Markel green, then yel-	Methylene green extra yellowish G 411. Turns greener.	XELLOW: Malachite green crystals or powder	Marachite green B. Benzal green New Victoria green New green New green Solid green Solid green Solid green Solid green	Solut green O. Vert soldo cristanx O. Diamond green B. Fast green	Brilliant green	Smaragd green Solid green J. Solid green J J O.	Victoria green 3 B	Fast green 248. [Yellow, excess water Fast green bittsh. 248. [greenish-yellow.	Methyl green Vert etincelle Domble green S F	Ashryl green 32g First green, then yel- Methyl green 32g First green, then yel- low and green 33k Brown-vellow

GREEN-Continued.

ont.isss—Confriend. New Victorin green. New green ergst. Solid green ergst. Solid green it was to blannon green. Vert collide erlekaux O. Fakt green.	Hellinat green Maheeline green (1 Maheeline green (1 Makayi gr	Solid green J J O	Light green S F yellowich	Christophia (green green	Frac green extru	Methyl green. Vert éthoetle. Lighti green. Lighti green.	Ashyl green
	2.100	Solid green J Solid green J Indict green J Indict green J Acid green J Acid green J Acid green D Acid green D		ppt.			_
Violet Continued. Signature Signatur	Metajaran green exan yenowasi et (44,) viore sai, mat viore content of the conte	Multachite green G Diamond green G Zibid green G Zibid green G Zibid green G Zibid green G Solid green G Solid green G	Guitnen green, B. 288. Mincklab, green 1914. Corruteline in pasto. 347. Dirty green colora- Corruteline A in pasto. 347. Litan, Corruteline B in nuccion or in mate.	Cartilland B. W. In powder or in Aff8. Olive-green profile and analysis of the profile and a second profile and a	Calinali green. 425. (Violet, on excess blue- Solid green G. green ppt.	Alizarina green H	parametric green is a power \$284. Ppt. of color base. Borzal green \$

On warming violet,	Carmine red.	Ppt.	Almost no reaction,
RED: Wool green S338. On warming violet,	Alizarlne green S	Fust green M 435. Ppt.	Gambline G in pusite
Turus yellower.	Red yellow.	Clear yellow sel,	Green-yellow fluoreservence.
Yerlow: Usamine green B	New solid green is B and 8 H	Chambine It lu paste, 376, Clear yellow sol,	Gambline G in practe Gambline Y in practe Gambline Y in practe Abarce green J

YELLOW.

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	_	
1	ō.	

Viol.RT; Sun voline	Hode vlotet	BLUE-Continued.	Yellow, with blue du-
Metanil yellow		Chromine 6.	огенсенсе.
	74.	GINEKN: Jame golde N. Jame green. Fast yellow.	Hue-green,
Metanil extra		Alizarine yellow F S 272.	
Janue metanile brome 75.	é	Benzolavine No. 2	2
Curcumetine New yellow Chronine Jampine	80. Red-violet.	Leather yellow Xantihie Philadabida wilaw (i	пиотемсевоз.
Janua respetant an saven.	ıć		S73. \ Kreen fluorescence.
Arnica yellow	A Breed selection	Patent phosphing	
Kreachine G		Rheonine	Brown with green fluorescence.
Chrystening R	_	Christing	
A refer the years A. District of the A. Belliant, veltor V.	7. J. Red-violet.	Thlochromine Thlochromogene, 481. Frue yellow, with Thlochromogene.	Pale yellow, with green fluorescence.
Chrysophenhe		Anreoline	
Diantine gold yellow 256. Mekang yellow 0 271. Mekang yellow R 281. Illne-vlolet.	5. 1. 1. Hine-vloket.	Bhown : Alcohol yellow G	-
Mkado yellow. 13. Red to blue, danne sollele N	k. Red to blue. 9. Blue-green.	Wool yellow R. 40, Narordna 81,	Yellow-brown. Yellow-brown.
Orrelazoi yeliow Mekong yeliow R. 281, Bine-vlotek	 Violet-blue. Blue-violet. 	First, yellow R. Yellow W. Yellow W.	88. Yellow-brown,

	Red-yellow, Reddish-yellow,	Slight fluorescence.	Light yellow.	Green-yellow, with very strong green fluorescence,	Hed-vellow with	дтеен пногеясенсе.	Light yellow. Red-yellow.	Pale yellow, with greenish thores- cence,	Brown.yellow. Yellow, with blue flu- orescence.
YELLOW—Continued. Gold yellow Repending yellow Actin yellow R. B. Tropucoline O. Orninge R. F. R. R. E. Junne H. L. R. R. E. Junne H.	Diamond yellow R in paste	Uranine	Chrysoline	Benzoftavine No. 2.	Xantiline Philadelphia yellow G.	Leather brown Property Phosphine II, N. P. Patent phosphine Nankin in cake.	Alizarine yellow C in paate	Primulino Carrudolino Polyetromine Polyetromine Thiochromogene Aurealino Sulphine	Thioflavine S
Brown, with green fluorescence. Red-brown. Slightly brownish.			Pale yellow.	Gold-vellow.		Orange-vellow.	Drange-yellow. Orange yellow.	ro. 77. Brown-yellow,	
374. 384. 384.	35	\$ E	જ	ನ್ಟ್ ಪ	si	83 23			ž
BROVN—(buttnucd. Ribeonine	to i, O, II, cono	Auramine G	YELLOW: Aurantia. Kaiser vellow	Pieric acid Tarfrazio Nitropheniue	Thinzol yellow.	Pragne alizarine yellow G. Alizarine yellow R. Orange R.	Terra cotta I. Alizario sellow R W powder. Prague alizarine yellow R. Persian yellow.	Fract yellow G. S. Fract yellow G. S. Fract yellow G. S. Fract yellow greenish Acid yellow L. New yellow E.	Acme yellow Glaysoine Claysoine Claysoine Yellow T.

= 3

YELLOW-Continued.

120. Brown.red. 103. Ormge-red. 170. Ormge-red. 447. Bluish-red.	480, Blood-red.	.487. Yellow-red. 488, Yellow-red.		SO, solution.	107. Plocky reddish - grny	252. Ppt. 281. Hlack-brown ppt.	381. Ppt. 382. Gray-white ppt.	108. Bine-violet ppt.	1196, i Bluc-violet ppt., then greenish-yellow ppt.
Alkali yellow B. Alkali yellow B. Catton yellow G. Heligaland yellow F. Flavinduline G. Othermodus A.	Oxyphenine yenow Oxyphenine Good Chlerophoenine Gold Oxyphenine Gold Thiophoephine J	Chinoline yellaw alcohol soluble, 487. Yellow-red. Chinoline yellow	YELLOW	On addition of H,O to H,SO, solution.	Chrome yellow D. Anordani yellow II N. Anordani yellow. Anordani yellow. Anordani yellow. Anordani yellow.	Mordant yellow O. Hassian yellow. 252. Ppt. Mekong yellow R. 281. Illack-brown ppt.	WHITE: Alizarine yellow A in paste	Violett: Cotton yellow (1	
Red to orange.	5. Red to opingo,	Pure cherry-red.	Red to blue, Dark blood-red,		81, Fuchsing red.	82. Illue-red.	Orange-red.	Orange-red.	Sourlet-red.
5 : : : : : : : : : : : : : : : : : : :		2 ±	13.				107.	108.	E E E
OHANGE: Mikado gold yellow 2 G, 4 G, 6 G, 8 G, 1 lirest yellow 2 G, 3 G	RED Mikado gold-yellow 2 G, 4 G, 6 G, 8 G. Direct yellow 2 G, 3 G Direct yellow 2 G Direct yello	Ourennine S Sun yellow Direct yellow G Jamie direct.	Mikado yellow	Azo yellow M. Azo yellow M. Azo acid yellow	Indian yellow C. Indian jellow C. Janno Dellant. Hellanthine. Citronine	Brilliant yellow B. Hrilliant yellow Y. Yellow W. H. Curenmino	Ohrome yellow D. Anthrecome yellow B. N. Mordant yellow Mordant yellow O. Mordant yellow O. Mordant yellow O.	Fulling yellow Crumpsall yellow	Oriot yellow R

Bress:		Bunw N - Continued.	
Fast yellow (Flust plue pat.	Steel-blue ppt.	Rheonine	=
Kresotine yellow R	•	Chloramine yellow	Inorescence.
Chrysophenine 251. Fpt.	(Ppt.	Oxyphenine Coxyphenine gold.	Ochre-brown ppt,
GREN:		Untophosphine d	
Kresotine yellow 1t	(Hine ppt., turns then green, finally yel-	Chrondne G Grownish flocks.	Brownish flocks,
Diamine gold yellow		Yellow: Anrantía. Kalser yellow	2. Ppt.
Buows;			3. Light yellow.
3	. Yellow-brown ppt.	Mikada gold yellow 2 G, 4 G, 6 G, 8 G.	
Nexy yellow Sit	S6, Olive-brown,	Direct yellow 2 G, 4 G) Thrtrazine	Ppt.
Azo yellow M		Thiazol yellow 9.	Turnslighter, no ppt.
~	SI. Vellow-red, later yel-	-1-	10. Brown-yellow ppt.
	the second of the	Direct yellow G	
Relianthine		Alizarine yellow G G	27. Light yellow ppt.
Nurceine	Yellow - brown, on warming exempes	Prague alizarine yellow G 28.	Ppt.
Heligoland yellow 170, Chrysonaline d 195, Carbazol yellow 200,	_	Albarine yellow R	Brownish-yellow ppt.
Mekong yethov G 281, Mekong yethow R 281, 281, 281, 281, 281, 281, 281, 381, 381, 381, 381, 381, 381, 381, 3	. Ppt. Ruck-brown ppt.	8 9	Orange yellow ppt. Brownleh yellow unt.

YELLOW-Continued.

YELLOW - Continued.			YELLOW—Continued		
Fast yellow G. Past yellow G. Past yellow G. Past yellow G. Past yellow greenish.		77. Orange-yellow.	\$ 352. \\ 353.	Yellow sol. with slight fluorescence. Ppt.	with cence.
Acid yellow			Acridine yellow	l'pt. Yellowish sol.	, then
Azo yellow M. Azo ned yellow Indian yellow	. St.	SI. { Yellow-red sol., then { yellow-brown ppt.	Touther yellow Nanthino Philadelphin yellow G Lenther brown Settler brown Settler brown	orange ppr. Red-yellow.	
Janne brillant. Relianthino Citronino			Thosphine 11, N. F. Patent phosphine Nankin in cake.		
Aema yellow.			479	Turns yellow.	
Chrysgoline Yellow T.			Primuline	,	
Gold yellow. Reserved by State of State	83.	83. Reddish-yellow.	Polychronine Thlochromagene 1481. Ppt.	Ppt.	
Acad Jenow R S. Tropucoline O			Aureoline		
Jampe II.	_		\$ 25	Orange-yellow ppt. Plocky ppt,	ppt.
Oriol yellow. Cotton yellow III3. Brown-yellow ppt. Alkull yellow.	113.	Brown-yellow ppt.	annothne yetlow		
Diamond yellow R in pasto	=	Brownish-yellow ppt.	RED: Alcohol vellow G		
Diamond yellow G in paste, Yellow gelatinous	.116.	Yellow gelatinous	Anilline yellow.		
Auranluo Auranino O, I, II, cono. Fuodino O, I, II, cono. Safrantino coarlet.	, 38%. -38%.	282. Pale yellow.	Alcohol yollow R. 40, Reddish put. Metanil yellow Orango M.	Reddish sol., reddish ppt.	then
Auramine G	5	Light yellow.		74. Fuchshie red.	
Yellow coralline	.330.	Yellow sol. and dito ppt.	Jame G		

Ppt. [Flesh-colored gelati- nous ppt, and red- dish-violet water- soluble crystals.	Red-violet. Ppt.	Steel-blue ppt. Greenish ppt.	5. Brownish to yellow.11. Brown, later dito ppt.13. Brownish ppt.	81. Brownish, on excess deeper brown.	Ppt. Ppt. Ppt. Ppt. Plocks. Ppt. Flocks.
VIOLERT: Janue metanile bromé. 75. Ppt. Fast y gelow G. Fast y gelow G. Fast y gelow E. Fast y gelow E. New y gelow E. Janue resistant an awom	Yellow fast to soap. Brilliant yellow. BLUE:	Fanne solide N. Steel-blue ppt. (GREEN: Diamine yellow N. 239. Greenish ppt.		Azo yellow Azo yellow M. Azo yellow M. Azo ilavine. Azo ellavine Indian yellow G. Indian yellow G. Junne brillant. Helianthine the Citronius.	Gotton yellow G 168. Heligoland yellow 170 Chrysamine G 185 Carbazol yellow 200 Chrysamine R 220 Chrysophenine 251 Diamine gold yellow 255
RED—Continued. Azo yellow Azo yellow M	Brilliant yellow S Brilliant yellow R Yellow W.R Curramine	Fast yellow Sast yellow Sast yellow Yellow W Sast yellow Yellow Yellow Sast Yellow Y	Jame resistant au savon. Yellow fact to soap	YELLOW. On addition of HCl to Water Solution. BLACE:	Chrone yellow D Anthracene yellow B N Anthracene yellow B N Mordant yellow Mordan

YELLOW—Continued.

Ppt. Orango-yellow ppt. Ppt.	Brown-yellow ppt. Hownish - y e 1 l o w Gacke. Brownish - y e 1 l o w Hownish - y e 1 l o w	282. Dark-yellow, decom- poses on boiling. 390. Yellow sol., and	Flocks. Brown-yellow ppt. Ppt.	.480. Orange-yellow ppt.	Pyt.	Orange-yellow ppt, Orange-yellow ppt, Turns lighter,
YELLOW—Continued, Prague alzarine yellow G	Alkali yellow H	Auraunino O. I. II. conc. Auraunine O. I. II. conc. Fuchsino searlet Safuraine searlet Softwa corallino Yeltow corallino	Uranino.	Chlerandiae yellow Oxyphenine gold Oxyphenine gold Thinphosphine J Chlorophenine G	Primutino Carnetino Polyeironino Tribodramageno A mrealine Sulpino	Thiodiavino S
Ppt, Dark-brown ppt, Brownish flocks,	4. Precipitation of dini- tro a-mapthol.	Precipation of tetra- nitro-phenolphtha- leine,	Ppt. Brownish-red wilth orange-red fluores- cence.	2. Light yellow. 5. Brownish to yellow.	Turns lighter; no ppt.	8. Ppt.9. Red-yellow ppt.10. Brown-yellow ppt.
Bhow N—Continued. Mekong yellow G. Nekong yellow R. Chrouding G. ASI.		Aureline	Reuzoflavine No. 2. 272, Ppt. Reuzoflavine No. 2. Incompleted with the theorem Reacher Reuzoflavior Reacher Re	YELLOW: Aumith Kalser yellow Mikado gold yellow 2G, 4 G, 6 G, 8 G, 10	5	:

No Reaction — Continued Trapecoline O	-	YELLOW.	On Addition of NaOH to Water Solution.	Auramine O, I. II. Par. soluble in ether; Auramine conc. Puchsine score. Capt. Safranice scarlet. Safranice scarlet.	283.	Benzoflavine No. 2	Violett: Nitrophenine	Green: Uranine Transcelluc Fluorescelluc fluorescence fluorescence
40. Red crystals. 74. Fuchsine-red and ppt.	Flesh-colored gelnti- nous ppt., and red- dish violet water soluble crystals.	80. Fuchsine-red.	Violet-red.	Fuchslne red, Orange-red.	Brownish-red, with green fluorescence.			
Abohol yellow R. 40. Metanii yellow R. 60. Orange M. N. Tropusolin G. Junne G. Junne G.	E	Chronine S0. Citronine Jasmine	Brilliant yellow S. Brilliant yellow Y. Yellow W.R. Cureunlue.	Frat yellow 88. Fuerbline rec Yellow W	Rheonine374 { Brownish-red, with	No Reaction: Picric acid Tartrazine	Acme yellow G	Yellow T. Reserving Reserving Plant Reserving Plant Reserving Plant Reserving Plant Reserving Plant R. S.

YELLOW-Continued.

			HROWN - Computer,	
				Trown in a hinning
Attachine yettow A 10 parte	381.	turns green rapid	Alizarine yellow C in paste,380.	darker on exposure
=	Ξ	ly.	Chilodavine in paste,	Yellow-brown.
4	=	Ornuge-brown,	ORANGE:	
			Mikado yellow	Yellow to red-orange ppt.
Oltronine	Z	86. Yellow-brown,	Prague alfancine yellow G	Dark red orange.
Azo yellow			Cotton yellow G	Deeper orange.
Azo flavino			Chrysopheuine	Yetlow set, and orange flocks.
_~	ž	81. Yellow-brown.	Мітова481,	Red-orange ppb.
Janes brillant			Yellow;	
Heilauthine			Auranine	2. Deep orange-yellow.
A ome yellow				3. Dark yellow.
Chryseofne System T.			Mikado gold yellow 2 (l, 4 (l, 6 (l, 8 (l) 8 (l) 8 (l) 9 (l)	5. Ppt.
	<u> </u>	83, Reddish-brown.		6. Orange-yellow docky
Jaune II Orange R L. Orange R R L.				9. Red yellow sol. and dito ppt.
Arnies yetlow.	2	Red-brown.	Cureumine B	10. Red-yellow ppt.
Chrysamine B	22(i.	Red-brown, Red-brown, Red-brown,	=	Ppt.
Ourysoline Ricceine 374.	37.4	Turns darker. Light brown put.	Alizarine yellow G G	27. Orange-yellow.

Turns redder.	34. Blood-red.	-	violet.red. Brown-red.	. Orange-red ppt.	. Turns redder.	Orange-red ppt. Turns redder.	Yellow-red. Cherry-red. Turus redder.	f Alcoholic sol. cherry- f red.	
RED: Tartrazine		Brilliant yellow S	:	Chrome yellow D	Oriol yellow R. Cotton yellow R. Albeit yellow R.	Alkali yellow R. 120, Helgoland yellow (V. V. V	Arcsonie yenny G. 250. Besian yellow 253. Mekong yellow R. 253.	Aurine	Orange M. N. Tropeoline G. Victoria yellow (O double conc.). Métanil extra.
	Green-yellow, Orange-yellow, Reddish-yellow ppt. Yellow sol., and	Orange nocks. Orange-yellow flocks. Orange-yellow.	Turns darker with dark-green fluores-cence.	Ppt. Light vellow ppt,	Deep yellow, turning	rapidly green onex- posure to air. Brown-yellow.	Yellowish ppt., solu- ble in ether without fluorescence.	Orange-yellow ppt.	Green-yellow flocks. Turns darker.
Yellow—Continued. Jame métanlle bromé	-	Diamine gold yellow S55. Alizarine yellow F S.		Acridine yellow	Phosphine II, N, P. Patent phosphine Narkin in cake.	Alizarine yellow A in paste	~~	Chlorantine yellow Oxyphenine goll Oxyphenine goll Oxphenine goll Chlorophenine G	Chromine G

YELLOW - Continued.

	Almost no reaction.		
Louis Standard Montage of CN	Vellow fast to soute. Yellow ourilline	Thiothering T. Thiothering 272, Thiothering T. Thio	Thiotheningono, Antwolline Sulphine Thiothwise S. 483.
Construct on .		New yellow 1. Itellihart yellow S	violat-red.
			7
	final yellow forst yellow G. S. Andry yellow greenish.	<u> </u>	- 4

ORANGE. On H₂SO₁.

Blue: Direct orange 2 R	15 blue 15	Colonities: Acridine orange	Orange G	Yellow: Cerotine orange C extra	Ponceau 4 G B. Crootine ornuge. 22, Orange-rellow. Pyrotine ornuge G. Pyrotine ornuge E N L. Crootine ornuge E N L. Crootine ornuge.	Orange G	Orange III (No. 3)
lue-gray.				Yellow-brown.			Violet-blue. Red-violet.
Black: Toluylene orange R R342. Bluegray.	Helianthine Helianthine Orange III. Orange No. 3 Directly bandline on ange No. 478.	Tropeofine D Gold orange Toluyene cange R. Kanthosine R.	Alizarine orange A in paste. Alizarine O R.	5389.		Tropaoline O O F. Past yellow Diphenyl orange Diphenyl orange Daylow Jawa d'antiline orange	H 8 8

ORANGE-Continued.

ORANGE.	On addition of H2O to H2SO, solution,	Иглеск : Аzo otrungo 1k	BROWN: Mikado orange G, R. 2 R, 3 R, 4 R, 5 R., 13. Hrownish ppt. Forecant 4 G B. Chredie orange Relitions orange G.	Orange G R X. Pyrotine orange Orange B N L. Orange B N L.	Orange G T, R N Brilliant orange O Orange N	Orange I Saphaladorange Saphaladorange Saphaladorange Saphaladorange Tropwoline O O O No. I Saphaladorange Saph	Orange R. Crange R. St. Yetlow-brown tlocks. Kernashie orange.	Orange R. 91 Ppt. Chicup orange	Tolin/done orange G { 221. Brownish ppt.	Ormige IV Ormige N (48 S. New yellow Acid yellow Acid yellow Yellow
Changes from red to	Jerum.	23. Cherry-red. j		84. Vloletred.		85. Fuchshæred.		S9. Fuchano-red,	Cherry-red.	
HED: Mindo orange G, R, 2 R, 3 R, 4 R, 13 Changes from red to	, de la constante de la consta	Ponceau 2 G	Orange (T. R. N. Orange (T. R. N. Hillilliant orange (O. Orange (N. C. S. Orange (N. C. S.	Orange I Niphthol orange.	Orange II.	Orningo A. d' Aelsl orninge Alliss orninge Gold orninge (36, p.	Mandarine U extra	Orange T. Orange R. Kermesine orange.	Orange R	Toluylene ornuge (1

Yellow - Continued. Alizarine orange A O P. Alizarine brown	RED: Cerotine Orange C extra	Helianthin Orange III Dimethylaniline orange Methyl orange Troppeoline D Gold common	Cotton orange G	Tolnylene ordinge R	Acridine orange	No Reaction: Ornige G. Ornige G G. Ornige G G en cristaux. Patent orange .	ORANGE. On Addition of HCl to Water Solution.	Видек : Azo orange R	Brown: Mikado orange G, R, 2 R, 3 R, 4 R, $\}$ 13. Brownish ppt, 5 R.
79. Ppt.	Denzo orange R	Orange III (No. 3)	Acrinine scarce 11, 2, 11, 3, 11	Yellow: Ponceau 2 G. Politint ponceau. Prillint ponceau. 23. Red yellow.	Orange III (No. 3) First crange relieve		Gold orange Mandarin G extra β-Naphthol orange Tropacoline O O O No. 2.	371.	389 Light yellow ppt

ORANGE-Continued.

 13. Brownish ppt. 190. Red-violet, 12. Turbid olive-green ppt. 191. So the in hirge suppt. 201. So the in hirge manner of verter of manner of verter 			.;; .;;	S5. Brown-yellow ppt.					400. Turns lighter, Orange-yellow ppt., so In ble in large nmount of water.			15.		78. Fuchsine red			
	VIOLET Continued. Fast yellow.	Diptemyt orange Diptemytamine orange Diptemytamine orange Tanne dynaline Renya orange 191 Brownish	GREEN:	Direct orange 2 K 12. Turbid olive-green Yellow:	Orange III (No. 3),	Orange II	Orange extra Orange A. G. Aeld orange	Atlas orange Gold orange	Mandarin G extra.	Tropicoline O O O No. 2,	OHANGE:	Orange III, (No. 3)	BED	Cerotine orange C extra,	Helianthine Orange III	Dimethyl anillne orange.	Tropucoting D
		 Yellow-brown ppt. Oily drops. 		13. Oily drops.	- F	- 1	Brown-yellow ppt.				89, УсПом-brown поскя.		Flocks, Ppt.	Yellow brown flocks,		Ppt.	
	_		_			-					_	_	192	221.		79.	
	Bhown—Continued. Ponceau 4 G B.	Croceme orange Coronage Brilliant orange Coronage Coronag	Orange E N L.	Orange G T, R N. Brilliant orange O	Orange I. Naphthol orange	a-Naphthol orange Tropeoline O O O No. 1	Orange II	Acid orange	Athe orange Red orange	B-Naphthol orange. Tropceolin O O O No. 2	Orange T.	Kermesin orange	Clileago orange	Toluylene orange G Yellow brown flocks. Kauthosine J	V1 DERT: Orange IV	Orange N, G S. New yellow.	Acid yellow D

NS. Dark brown.	Orange T. S9. Red-brown. Orange B. S9. Red-brown. Chicago orange	Mikado orange G, R, 2 R, 3 R, 4 R, 9 Yellow-to red-orange B Mikado orange G R, 2 R, 9 Mikado orange C extra Mikado orange for cotton Mikado orange for cotton or	22. Brown-yellow.	23. Turns yellower.	78. Crange - yellow ppt., soluble in large amount of water.	Orange R
Brown-red ppt. Orange II. Reddish ppt. Orange extra. Acid orange. Arboneoline O O O No. 2.	Orange T. Orange T. Kermeshie orange. Chicago orange.	red. Mikado orange G. II, 2 II, 3 IR, 4 IR, 6 IR. 4 IR	23. Almost no reaction. Ponceau 4 G B. Croerine orange. Frilliant orange Orange G R X. Pyrotine orange E N L. Orange E N L.	Joneana & G. Brilliant poneau G G. Ponteau J J. Helianthine.	vn.	I3. Dark brown-red. Orange R
RED - Continued. 91, Cotton remge R. Cotton remge Cotton rem	Acridine orange R extra S71 Orang Alizarine orange A in paste Orang Alizarine O H	Alizatine orange A O Alizatine orange powder. Alizatine orange powder. Alizatine brown	No Reaction	Patent orangeORANGE.	i	Orange G T, RN. Brilliant orange O. Orange N.

ORANGE-Continued.

.1. Sl. Cherry-red.	Cloth orange 192 Turns red, in conc. Tolylene orange R R Paramine Paramine R R Paramine R Paramine R R Paramine R R Paramine R R Paramine R R	- SS9. ← Fr	Alizarine orange A O P	111.	201,
lish orange-yel-		Alizarine O B. Alizarine O G. Alizarine orange N. Alizarine orange N. Alizarine orange A O. Alizarine orange A O.	Alizarine orange A O P Alizarine brown	No Reaction; Cotton orange G	7.9. Egg-yellow ppt. Toluylene orange R Kanthosine R. Gongo orange R 215.
YELLOW—Continued. Tolaylene orange G Kanthosine J Acridine orange Acridine scarlet, R, 2 R, 3 R Acridine scarlet, R, 2 R, 3 R	Obligate orange 2 R		Orange G T. Orange R N. Brilliant orange O. Orange N.	Orange IV Orange N Orange G Orange G Now wellow	Acid yellow D. Tropacoline O O. Fast yellow we. Diphenyl orange. Diphenylamine orange.

BROWN. On H_2 SO.

Black :	14. f Red black to blue- black. Violet-black. 277. Violet-black.	Violet—Continued. Alkali brown R. 110. Blue-violet Beliac brown B. 170. Blue-violet Terra colta F. 161. Real violet 161. Real	110. Blue-violet.
Brown: 87. Red-brown. Chrysodine R. 11. Red-brown. Leather brown. 160.	 Red-brown, Red-brown, 	Fast brown G. Acid brown Fast brown (O N T yellowish). 165.	
Account nown Bismarck brown Bismarck brown G Manchester brown Phenylene brown	165.	Diamine brown 7 183. Cloth brown reddish, 1133. Cloth brown yellowish, 194. Diamine bronze G	Blue-violet. Blue-violet. Red-violet. Blue-violet. Red-violet.
Anilhe brown. Leather brown. Chunamon brown.	172,	Congo brown 14. 270. Benzo brown B. 274. Toluylene brown. 276.	Dirty violet. Dirty red-violet.
Canelle Buglish brown		Blue : Pigment brown	
\sim	174. Dark brown.	Fast brown 3 B103.	
Bismarck brown T. Brun Blsmurck E E. Benco brown G.	773. Violet-brown.	GREEN: Sulfamine brown A	
	65,	Yellow: Chrysoldine G. Chrysoldine G. Chrysoldine R, J, Y.	16. Brown-yellow.
Naphthylamine brown Fast brown N. Azo brown O. Clurome brown R O.	÷.	RED: Fact brown. 164. Corinth-red. Toluylene brown G. 241. Brown-red.	Corinth-red. Brown-red.

BROWN - Continued.

Harawa Continued, Terra cotta F., Hasareta braya, Fall Hall Proven Fall braya, Hall Hall December of the Cotta Cotta	First Inner O. Acid Inner D. Yellow Brown.	Noncylne II Manchester brown B. E. P. S. Hamarel drawn. Hamarek frown T.	189	Cleft brown yellowith, 1914 (1944) Congo brown (1) Congo brown (1) Congo brown (1) Congo brown (1)	-	Henzo brown H 274, Pul. Pul.	_	Allanding brown powder Visition W. Iron P. M.	Hen :	Ohrysofilline R
 Charify rad.		THOUGH TOUT	Brown red,		olntion	Hue black ppt.	Ppt.	Ē Ē	Yellow Igowu,	Durk brown and
Heb Continued Palyobrantine H Colton brown R Finst colton brown R	Authinema brown in pacto, Anthracema brown Hitpacto , Anthracema brown O in pacto ,	Altarine brown in paste. Altarine brown pawder. Kutgallel	Thiographical Thiographical Thiographical Thiographical Thiographical St.	BROWN,	On addition of H ₂ O to H ₂ SO, Solution	yetromina B. Iton brown R. act collon hrown R. rede frown R.		I 15 1	Submittee brown 16. 18. 185.	Alkall brown R

Ppt.			Turns lighter. Ppt. Ppt. Red-brown ppt.		Chocolate-brown ppt. Ppt. Ppt. Ppt. Ppt. Ppt.	Ppt. Ppt. Ppt. Dark brown ppt.	rpt. Ppt. Ppt.	Red-violet.	Fpr., soluble in dir. HCl with violet, in water with brown color. Ppr.
BROWN: BROWN: BRING DOWN B, 3 G O, M. 14.	Chrysoidiae R. 41. Alkali brown R. 110.	Benzo brown 5 R	Leather brown 160 Terta cotta F 161 Reacrin brown 163 Fast brown 164	Vesuvine B. Manchester brown E. 174. Manchester brown P. 174. 174. Bismarck brown P. 174. 174. Brum Bismarck brown T. 174. 174.	Diamine brown V 182	Congo brown R Z70. Benzo brown G 273. Benzo brown B 273. Direct brown, J 275.	Toluylene brown B. Hessian brown B. Hessian brown M.M	VIOLET: Fast brown 3 B103. Red-violet.	Fast brown G. [165. High Acid brown (O N T yellowish)166. Ppt.
41. Slimy ppt.	112. Fuchsine red.	.99		> 172.	.174. First red, then brown.	92. Brown-red ppt.		er Solution.	13. Blue-black ppt.
IRD—Continued. Chrysoldine R	Naphthylamine brown Fact brown N. Aroo brown O. Chrome brown R O.	Fast brown (O N T yellowish)166.	Bismarck brown G Illiamarck brown G Manchester brown Phenylene blown	Antline brown Letther brown Clunamon brown Canelle English brown Gold brown	Vesuvine B. Manchester brown E.E. Manchester brown P.S. Bismarck brown P.S. Bismarck brown T. Benn Blemarck E.	Thiocatechine S	BROWN.	On addition of HCl to Water Solution.	Flower Physics Property Property Property Property Past cotton brown R. Direct brown R.

BROWN-Continued.

uea.	BROWN.	On Addition of NaOH to Water Solution.		Ohrysoffilme C	Chrysofiline It.	Chrysofidine Y	Brun P M 97. Yellow-brown ppt.	Sulfamine brown A	Sulfamine brown B. (65. Yellow-brown ppt. Narhtdine B.	Naphthylamine brown	Ago hrown 0. College of the prown of the prown of the prown of the prown it to	Leather brown O 160. Ppt.		Blemarck brown G	Phenylene brown	_	Leather brown	Chunamon brown	Canetie	Gold brown	Vемиуine H	Manchester brown E E.	Manchester brown 1' S 17.1. Light-brown ppt.	Hismarck brown 7	Run Hanarck E. E.
DIOW IN - Comeruned.			FIG. INOMI SCHOW HOCKS. Huo				92. Fuchshie red.								651										
	YELLOW:	Chrysoftline C	Chrysoldine J.	Chrysoldine Y	1660:	Naphthylamine brown	Aza brawn 0	Chrome brown It O.	No leading:	Brun naphtine a	Sulfamino brown B	Bismarck brown	Brun Blannrck	Manchester brown	Venvine	Annue brown	Chnamon brown	Canelle	Gold brown	Anthracene brown in maste	Anthrucene brown A in passic	Anthracene brown G in pusto, Mill.	Allzurine brown in paste	Augustine mown powder,	Rufigallot 408.

Fuchsine red.	Ppt.	Cherry-red. Cherry-red.	Door red	Almost no reaction.		
RED: Fast brown 3 B	Alkali brown it Benzo brown 5 it.	Fast brown Fast brown G. Sherry-red.	Congo brown G. 269, Congo brown B. R. 277, Done and Heeding brown B. R. 977.	No Reaction: Mikado brown B, 3 G O, M	Toluylene brown G. 275. Toluylene brown J. 275.	
Ppt.	In conc. 501. small quantity of ppt. Red-brown.	Ppt.	In cone. sol. flocky ppt. Red-brown.	Green-blue.	Unstable.	Ppt. Red yellow. Turns yellower.
Bnown - (batimurd. Diamine brown V182. Ppt.	Cloth brown reddish	Polychromine B. Cotton brown R. Fret cetton brown R. Briest cetton brown R. Direct brown R.	Benzo brown G	BLUE: Anthracene brown in paste Anthracene brown it in paste Anthracene brown it in paste	Alizarine brown powaer) Ikufigalloi	Verloow: Chrysoldine R Chrysoldine R Ppt. Pest brown (O N T yellowish)166. Red yellow. Diamine bronze G

RED.

172, Gray-violet. Hue violet, Blue-violet. Hine-violet. Hine-violet, Red violet. Red-violet. 125. 1125 Procedue B, A Z, 124. tessinn purple H. 247. drythrine X..... tensian Berdeaux....... Noth red O, E. Azo Corlinth Joffan semlet, Allzarino Bordenux II in paste 20% Mugdala red) Rouge L..... Hamilto scarlet, 3 B. Manilue Bordeaux II... Alizarine Bordeaux G Chamber kase magdata, trilliant red... tubidine,.......... Jernsthe trilliant croceins Willlant croceins bluish fotton semilet a B conc. 'aper searlet blutsh..... Alizarine Berdeattx B D m paste Alizarino hordentix G...... williant crowine M...... 'onceau B O extra...... Applicht Jamine plate, Violet's Continued. Bark vlolet. Dark violet. Hed-violet. Red-violet, ₹. 5 = Ξ 'unt red B. 101. Chromotrope 2 B..... Azo Turkey red..... Fast red H T Azo fueliaine G. Crystal peneran filk. Oryntal poncenti... Azo ruby A. B. Azo fuebsine aelde tougo rubba A Poulpre Vaplithol red 8, 0, Closoria mby Weel red extra..... Enanthining fast red A fast rod tf. Carntolalne Fast red D, E B, N S Azor roby Brilliant carmolalne O... Azn acid ruby Amarantb...... Azo neid ruby 2 II. ... First rod C... Azo fuetigine II.... Actd carmolaine II..... Bordeanx D B, S. Peneran ti R. Roxamine RAL POL

Dark blue			181. 191. Red-blue.		
ULUE — Continued. Croceine scarlet O extra. 141. Fast pouceau 2 B. 142. Orselline B B. 144. Bordeaux G. 145. Croceine scarlet 7 B. 145.	Croecine searlet 8 B 146 Ponceau 6 R B 150 Brilliant eroceine 9 B 156 Glycine Corinth 176 Glycine red 156 Glycine red 177 Glycine red 17	178. 178. 178. 178. 178. 179.	Rougo M	Definition of the properties	Rosazurine G
Gray-vlolet.			Green-blue. Dark green-blue. Dull green-blue.	Binck-blue.	Dark bluc.
Violekt—Continued. Fast plak for silk Naphthalinered Naphthaline plak Naphthaline warlet Soudan red.	BLUE: Aso Bordenux Burdio ruby Palatine red Aso red A. Nanitho ruby Sanitho ruby Sanitho ruby Sanitho ruby	Fast red B. Bordenax B. G. Bordenax B. A. Ceradine Rouge B.	Chromotrope 10 B. 63. Congo G R. 119. Cloth red 3 G extra. 726. Cloth red 3 8 G A. 127.	Cloth red 11. 129. Croceine 3 B. 130. Cloth red 6 c.	Fast Stordenux O. Azo fuchkline acido Oreellis root A. Union fast cinret Fast scarlet B. Croseline scarlet B. Croseline scarlet B. Fonceau 4 B. B. Fonceau 4 B. B. Fonceau 4 B. B. Fonceau 5 Fonceau 5

		illd, Yellow mown,	346, With evolution of [HCl, yellow-brown,		338. Red-brown,	
<u>=</u>				- 18 P		# #
Guera Continued, Rafrantine G O R, R. Rafrantine G H PU, R. Rafrantine P F extra No. G. Rafrantine P F extra. Rafrantine A G extra. Antiline pith. Full Full Full Full Full Full Full Full	Huown i Fueliation Ruper the inagents eryshis flest inagents Ruby magents	Ulaurant groa erfatoux. Amiliae red. Nitrobergaal frebiabus Hoomilijne base	Bhedamine II. Historianine J. extra. Historianine J. extra II. Safraniline	Rose Bengale 314. Rose Bengale 314. Altantine granufe H. Altantine granufust.	Albachie R G and G L Albachie S D G Albachie S D G Albachie O D Albachie O A Flavorie O A	Contouring red
	Who green,			DAIN ETCON,	Bluty green, Blety green,	
Chaline red N O. Dientine red N O. Dientine red N O. Rouse preparation in H. Naphthy Jene red Resigna juryle N Hessian turyle N Hessian fulliant juryle WHEN I	Obrasilia red (19), final G. (19), f	Foursont Bestra. Vest poursont B. Sowred L. Storriot B. Storriot B. C. Integral mariet.		= = =	Londontine II. Rodinduline 2 II think Rodinduline 2 II think Rodinduline 9 II think April 100 III think April	Safranie T. A. Safranie Stra G. Safranie Stra G. Safranie G extra

YeLlow—Continued. Easine alcohol soluble Prinnes & Falcool D H Methyl cosine.	Bowlin B, B B. Rose J B i Pateool. Alcohol cosine Ecsine atcohol soluble. Frimres a l'atcool. Primres a l'atcool.	Bowline B.N. Methyle ossine Safrosine Safrosine Safrosine Bosine scarriet B. Ecarlate J. J. J. V. Nopuline Bosine scarriet B B oxtra.		John ooking G	Jodine cosine B Rose B 4 Feature 2554 Perotuck 2564 Perotuck 2564 Perotuck 2564 Perotuck 2664 Perotu	Phloxine P. Phloxine B. Ergthrodine H.B. New pink.
18. Brown-yellow.		Brown-yellow.	Red-yellow, Red-yellow, Yellow color and green fliorescence. No fluorescence,	Greenish-yellow, Brown yellow.		
Ceratine orange (f	New fuchaline Leo Ruity Frochaine 8. Acid magenta Ruity 8.	Anton months and a second	Pyronine G	Anisoline Anisoline Anisoline Arisoline Arisol	Eowine A Eowine Pellowish Eowine gellowish Eowine G extra. Eowine G G F. Eowine water solutific.	Fosine 4.3 extra Eosine extra Kosine K. Sord. Eosine D. H., J. J. F.

RED - Continued.

	Poncoau G. Poncoau G.R. Poncoau G. Pelliant poncoau G.		rry-red. Agardne 8		Cochemille red A. Cochemille red A. Cochemille red A. E. Cochemille red A. E. Cochemille averled.	Industrial Process of
On warming, brome evolution, or brange yellow, on warming ford in evolution,	-yellow.	20, Fuchsine red.	Sherry-red, knby-red, Fuchshe red.		Fuchsine red. Ruby.red. Fuchsine red.	Fuebsine red. Puebsine red.
362, Ore 362, Ore 362, Week	1117, Ned	2d, Fue	25.	30.		
	Operation biot. Resident 10 II. Purpurine in paste. Alizardie No. fl.	RRD: Southen I. Carmine naphtha	Gochenille warlet G. Chrometrape 2 R. Oracillo annellinio V	Oracille aubatinte 3 V N.	Apollo red Oracille aubstitute N extra. Chromatrupe Ap. Cochonille search 2 II.	Red B Sondan II.

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RED-Continued.

On addition of H2O to H2SO, Solution.

KED.	On addition of H ₂ O to H ₂ SO ₁ Solution.	BLACK: Brilliant Purpurine R209. Blue-black ppt. Brilliant congo R	Distriction of the control of the co	Hessian brillant purple	VIOLET: Chromotrone [0 R. 63. Red-vlolet.	Fast red D, E B, N S. Amuranth Associate ruby 2 B.	5	Voltred extra	Crocefine B, A Z, 123. Red-violet ppt,	Crocofine scalet TB, 8 B. (146. Red violet. Poncoun 6 R. R.	Glycine Corinthe	Gradialine O. 1. 2	Roszurine G. 207. Red-violet ppt. Roszurine B. 208. Ppt.	
Double granter out on the	Double Exister Exist B. Bomble Orllant genetet 3 R. Exartate brillante Frillant nonceau 4 R.	Double scarlet 2 R. Fast acid ponceau Ponceau acide.	Fast acide scarlet.) Salmon red 169 Fuchsine red. Salmon red 171. Fuchsine red.	Cotton ponceau. Cinnabar scarlet. Since Fred.	Authracene red	Ronge de St. Denis D'anten rock scarfet 239.	Acld and milling scarlet	Alizarino V 1. Alizarine No. 1. Alizarine R. Alizarine So. 1 B new \$88. Deep yellow-red.	Alizarine for violet		 Alizarine S X.	Alizarine G D	Authra purpurine	

RED-Continued.

Hu. First blue sol., then brown-red flocks.	Blue to red. First blue sol, then brown-red ppt. Violet-blue ppt.	Ppt. Ppt.	Ppt. Violet-blue ppt. Later fuchelne red.	Later red.	۰
BBUS—Continued. Ponceur 3 B SATE Foureaur 4 B B Foureaur 5 E SATE First ponceur 1 B Sarrie B C Old scarle B C Old scarle B Imperial scarled. Establish SATE Blebrich scarlet.	Cloth searlet R	178,	Remo purpurine 10 ft. 1341. Ppt. Hessian fordeaux. 244. Violet-blue ppt. Noutral red. Rougo neutre extra.	Safranino T. Safranino Stra G. Safranino extra G. Safranino G extra. Safranino G 4 8 8. Safranino G 10 8. Safranino G 10 8. Safranino G 10 9.	Sufrantine cono. Sufrantine A G extra. Sufrantine A G T extra. Plink
		Mi. Vlolet blue,	Ppt. Later red ppt.	First blue sol., later brown ppt.	Later red sol. Then red to searlet-
Violent — Confinued. Congo 4 R. Rouge Congo 4 IL. Heasian Bordoux 244. Illuo-violet ppt. Rose ungfula. (Magdala red.) Napittylamino pink Fast pitk for elik Napittylamino pink Napittylamino pink Napittylamino pink Napittylamino pink	Napithaline scarlet. Sondan red. Blux: Fast red D, E B, N S. Amaranti.		Generatibiline) Occups of R. 119. Ppt. Southan II. 1 Ppt. Certaine red 121. Later red ppt.	Brilliant croceine M	Ponewat 5 R

					,) I E	A	٠١.	ALL	515.						-	219
	First bine schillen.		Red-brown flocks,	Yellow-brown p p t., with more water red sol.			Ppt.	Dark brown flocks.	Ppt.	Yellow-brown ppt.	Ppt.	Reddish-brown ppt.	Hed-brown sol., luter red flocks.	(Slightly brownish	and the	Red-brown ppt.	Red-brown ppt.
_			13H.	137.	ž		J. IBI.	201	1 2015.	₹ 20%.	2.17	27.	.380.	-	, JOK	361.	, tar.
Brilliant croceine M	Brilliant crocedne bluish Gottom scarlet. Cottom scarlet 3 B conc.	Poncenu B O extra	Oracilla red A. Union fast claret	Groceine searlet 3 B	Diamine scarlet B	Diamine Bordenux B	Diamine fast red F	Benzo purpurine B	Diamino red B	Diamine red 3 B	Hessian purple B	Azo Corinth	Acid rosamino A	Phloxine P.	Erythrosine B B. New pink.	Cynnosino alcohol soluble,, 361. Red-brown ppt	Parpurine in paste
Turning red later.			300, Almost colorioss,			Light brown ppt.	50, Yellow-brown ppt.		59. Ppt.	Red-brown ppt.	Yellow-brown ppt.			Yellow-brown ppt.			Ppt.
.50	181		.jog			. 18.		_		- 66.	.101			20		_ ~	- 1r6.
JEERN: Induline scarlet	OLOBREMS: Rouge M. 181, P-Fuchsine. 249.	Fuchsine, Ruby, Magenta, Roseine,	Diamant gros cristaux	Romaniline base	· wwo as	Cerusino orange G	Paintine searlet	Cochenille scarlet P S	Palatino red. Azo red A. Naphtho ruby	Azarino B. Red-brown ppt.	Fast red Bput. Yellow-brown ppt.	Fast red A. Frat red	Fast red O. Roccolline.	Kouge 1 Brillandr red Rubidine	Rauracienne	Fast acid pencent.	Poneran neide Parlet Ppt.

RED - Continued.

		Visitow - Continued.
O. Oral	20. Orange yellow ppf.	Pyrodine 14, 15, 15, 16, 16, 16, 16, 16, 16, 16, 16, 16, 16
-Hi, Stra	Straw-yellow ppt.	Kone B A Penti. Printone achible. Kentoo bin'eta.
2, Ited	52, Red-yellow.	Scaline denderse, Dhardrine B. Alizarine V. I. No. 1, Le.
fulling orange	Gray-yellow ppt.	Albarine No. f. Haew.
	Mob. Brown-yellow lpt.	Alizarina rod's. Alizarina W R powder W. A Alizarina operer W, S A. Alizarina operer W, S A.
E	e nist, - Brown-yotlow ppt,	Alizarine B G and G I, S B G, X, No. 14, O A. No. 19, O A. No. 19, O A. No. 19, O A. Alizarine S X, S X extra, G D, B X, S G Anthra purpurine Anthra purpurine Anthra purpurine Anthra purpurine Anthra purpurine
		Rowlinding 2 G
	Brown-yellow ppt.	Chromothip scarlet G
d	Ppt.	Chromotrope p B

DYE ANALYSIS.

98. Fuchsine red.	Brown-red ppt.	Yellow-red.	Brown-red ppt. First blue, then red.	Dark brown-red ppt.	Dirty red. Ppt. Violet ppt., then fuchsine-red sol. Brown-red ppt.	132. Brown-red ppt. 133. Brown-red ppt. 135. First blue, then red, then searlet.	Yellow - brown ppt., with more water red sol,	First blue sol., then brown-red flocks.
RED-Continued. Ponceau 6 R. 98	Double brilliant scarlet G	Double scarlet extra S., Double brillant scarlet : IR, Beralate brillante. Brilliant poncean 4 B. Double scarlet 2 R.	Cloth red G. R. [129] First red T. B. Ponceau 5 R. Brythrine X. [125]	Cloth red 3 G extra	Cloth red 3 B extra	Cloth red B, O, B A Fast Brodeaux O. A 139.	Fast scarlet B	Ponceau 3 H B. B extra Fact ponceau B
Reddish ppt,	Brown-red ppt.	58. Fuchsine red. 60. Dark brown-red ppt.	61. Fuchsine red.	Scarlet-red.	Yellow-red, Brown-red ppt, Yellow-red, Ppt. Bluish-red.	93. Fucirsine red.	Yellow-red.	Yellow-red,
ED—Continued. Cochenille scarlet 4 R 48. Reddish ppt.	Wool scarlet R. 49. Brilliant orange R. Scarlet G R. R. 51.	Ornige N. Azo Bordeaux. J. Buffalo ruby. S. Fast red B T. Fast red B T. 60.		Crystal poncean 6 R	Azo turkey red. 64. Azo cosine. 47. Azo exhentile. 68. Erika eb. 72. Azo fuchsine G. 57.		Acid carnoisine B 94 Croceine 3 B X 95.	Cochenille red A. Cocenille awarled B X. Brilliant sorbed Polineau 5 R. Ponceau 4 R. Ponceau brilliant 5 R. Sowyen earchie

RED = Continued.

Drawline availet O extra 141, Noveline availet O extra 142, O excline to U. 143, O excline to U. 143, I horizona to U. 143, I horizo	Veltow real, Yellowereal Fuchantur real, Pro- Huline subarbon, then Huline real, Hunder real, Pro- Pro-		
1147, 1764 1189, 258, 258,	the solution, then brown-rod ppt, fush red, fown red ppt, 'pt,		
, 25H,		Links of the same	dåt, Voltowered ppt.
Soll,	- 14	Control of the ord.	
		Bone Beignie N. A T. G	Brown red flocks.
And and milling nearlot, , 2019, 14	., 240, Sentletined,	Phloving	
New Fuchshine,	dath red.	Phiathe latin free, 1944.	Vellow-red flocks.
Aeld magenta,		Unantality H. Hone Trouble 3 H. Hill.	Pleath cudored ducks.
310	Oraqually reddendug.	Allearing granut R. Allearing (300), Carading red ppl.	Carmine red ppt.
Greint B		-	JUN, Hrownered ppt.
	Turns red. Turns red. Ornuge, then red.	Noutral red	449. Fibrat blue, then fuch atte.
	Pluk.	Induline sourlet	Phut green, then red.
Rhodanline B, B extra	earlet red into blus-	Rosinting 9 tt	Ppt, First yollow, then or- ange red,
Aplediating 3 B		Safrantine Safrantine T	dell. Piret bine, then red.
Acid Rosamine A (2001. Red-hywyn sels, llien Violentine G	red brown sol., Hien red flocks.	Safraulne G extra	

RED.—Continued. Sufrantine G U U U. Sufrantine F F extra No. U. Sufrantine One. Sufrantine A G extra. Antiliue pink.	. First bine, then red.	VIOLET	Ppt. Blue-violet. Ppt. Ppt. Ited-violet.
Wose magdala (Magdala red)	Violetred ppt.	Salmon red	Hine-violet ppt. Hine-violet ppt. Ppt. Brown-violet ppt. Red-violet ppt. Red-violet ppt. Ppt.
REACTION Applier to Substitute N extrem. Proceedings also that A R. 53. 63. 64. 64. 64. 65	Solution	Dinmine red N O 225, Naphthylene red 240, Sufranine T 240, Safranine T 240, Safranine Satur, G 25, Safranine G 0 0 0 G 25, Safranine G 0 0 0 G 240, Safranine G 0 0 0 G 240, Safranine cone 240, Safranine A G 24, Safranine A G 24, Safranine A G 24, Safranine A G 24, Safranine Dink 27, Safranine Dink 27,	Ppt. Black-violet ppt. Blue-violet.
CK: Brilliant purparine B 209, Najbhykien red 24, Hessian purple N 21, Hessian purple B 24, Hessian purple B 24,	Ppt. Ppt. Violet-linek ppt. Blue-black ppt. Blue-black ppt. Brown-black ppt.	Rose magdala (Magdala red) Naphthylmine pink Fast pink for silk Naphthaline red Naphthaline pink Naphthaline pink Naphthaline pink Saddan red	-472. More violet.

RED. - Continued.

51, Brownfaltred ppt.	M. Hrown ppt.	60. Dark red-brown ppt. 62. On exerne brown 62. Crystals ppt. 67. Ppt.	fig. Stimy frown sub- staties.	II. Yellaw-brown ppt.		102, Yellow-trown ppt.	104. Ррб.	ib. Yellow-brown flocks.
	Palating red Azorest Azorest A Napiting ruby.	First red B T. 690. Crystal parceau B E. 602. Crystal parceau B E. 602. Azo contine. 607.	Azo ruby A. 8. Azo andi ruby Azo andi ruby Azo furuhina nedde Frat red O Cermioline Livilland evrmulatio	Ronge ruby A fant. red II Fank	Fant red A. Rowelline	Rubilint ved Rubiline Eunvedenine Cermenen	Double brilliant agarlet G	Domble worket extra 8. Trouble Pellant wastel 3 R. Kewrine bellante. Upple worklet 4 R. Double worklet 4 R.
Hinish, Violet blue, Frite		Violet blue ppt. Violet blue ppt. 1'pt.	Ppt. Ppt. Ppt. Ppt. Ppt. Ppt.	excess pure fille,	-	With liftle acid green crystals, on excess nearlet-red sol., wilch turns bluist-red on diffution.	20. Brownish-red ppt. 41. Fartfally red brown	43, Referencewin proc. 50. { Telions - brown gela- for. } throns ppt.
Oracillo autoettuto 3 V N	Cloth red I II evtru, 127,	Sultron red 10th Sultron red 17th Congo 17th		Houge neutre extra, (****) excess pure blue, Garages :	Cloth red 3 15 extra 197.	Rhodaulne II	Oneille substitute V	Orabenille scarlet 4 R Pankline scarlet 2 R Brilliant cachenille 3 R Godsenille 4 R Godsenille scarlet F 8

Ppt.	Red-brown ppt. Brownish-black ppt. Ppt. Reddish-brown ppt.	Yellow-brown ppt.			Ppt.	In alcoholic sol.	appears.	364. Decolorized and red-		Turns yellower. Orange.yellow ppt.			Gray - yellow gelati- nous ppt.
Brows – t'ontinued. Diamine red 3 B. Delta purpurine 7 B. Rouge diamine 3 B.	Brilliant Congo R. 210. Hossian purple B. 247. Hessian purple D. 248. Aso cerinth. 27.	Eosine S. B B. Rose J B à 1 áteod. Alcohol Bosine. Rosine alcohol solitile. Rosine alcohol solitile.	Primrose à l'álcool	Alizarine granat B	Azo carinine B	COLORLESS: (In alcoholic sol. Cyanosine alcohol sol. 361. diorescence dis-	Phloxine	Flloxine T A Eosine bleu	Cyanosine Eosine 10 B	YELLOW: Chromotrope 2 B	Azarine R	Cloth searlet G	Fulling orange 138.
Ppt.		.124. Brown ppc.	Flocks,	Dark red-brown ppt.	large excess green-	132, Brownish.	Brownish-red ppt.	i Yellow-brown flocky ppt.	Dark red-brown ppt. Brownish.	Brownish-violet p; t.	Brownish,	Ppt.	Ppt-
Fast acid ponceau. (Ponceau acide).		Cotton scarlet. Cotton scarlet 3 B conc Ponceau B O extra. Paper scarlet bluish.	Ponceau 5 R	Cloth red 3 G extrn	Cloth red 3 B extra	Cloth red B. C. 3 A	Bordeaux B X	Groceine scarlet 3 B	Bordeaux B X	Brilliant Congo Θ R. Rouge Congo brilliante Θ Brownish-violet p.t. Gladioline Θ , 1, 2	Ronge M 18. Brownish.	Dlamine fast red F	Diamine red B '945. Ppt

RED-Continued.

Yerlow—Continuer, Egythrodine B, B, Egythrodine B, B, Torinne B, A Penu, Romerone achible, Ferincerone achible, Fordine blunk thinks Fordine blunkire Fordine blunkire	Priloxine P. Ergelrowine H B. New pink. Advanture red B.	Alizarine W 8 powder Y Silto. Light yollow, Alizarine powder W Alizarine powder B A. Alizarine condition on the silter of the si	Residuation 2 C	One North H. Azarlin H. Azarlin H. Azarlin H. Trenine yellow ppt. Ivenine H. Trenine H. Jahr Grange yellow ppt.		A pulle real Openition Newton (31. Puchaline real,	Octioning society It. 142. Friedle. An execute It. Double souther It. 17. Brown red fleeks.	Wool scarlet It 48, Blue red.
			On dlf, red, On dll, red,	355. Brown-yellow ppb.	Green-yellow fleeks,		358. Brown-yellow ppt.	
380	E.	.181	3146.		. 367.		358.	
Yealow — Continued. p-Fuchsine Fuchsine Fuchsine Superflue inagents crystals Heel inagents Ruly inagents Ruly inagents Ruly magents Ruseline Ruseline Ruseline Antiline red Antiline red Michenzale furthine	Romanilline base Magesita base Magesita base Silver fucibatine In rithy	Coralline Red coralline Aurino R Aurino R	Rhedenfor O, G extra. 1945. Rhedening 2 R. Ankelline. Rtf. Rtf.	Abadiol sol, costine Primrose à Libraol D H. Methyl costine Rostine B N, B, B W, D H V.	Mathy couline Material Material Material Manufaline Maphillian Maphillian Material Material	Kajeur red. Latarkeune	Dianthine G Pyrodine J Iodine codne G	Janua d'Orient

				pt.s			red ish-				-		
	Brown red.	Flocky ppt. Ppt.	Ppt.	Conc. acid red ppt., dil. acid crystula	On dilution red.		346, which turns bluish-	'un ou au	On dllution.	Flocks.	454. Yellow-red		Brown-red ppt.
	<u>z</u>	258.	259.	3:14.	345.	_	346.	_	347.	320			ğ
RED-Continued.	Diamine scarlet B. Diamine Scarlet B. Diamine Bordeaux II. Diamine Bordeaux S.	Authracene red. 198, Rock scarlet Y S. 258,	Rouge de St. Denis	Ithodamine 6 G3.14.	Rhodamine G. stra	Rhodamine B	Ithodamine H extra	Safraniline	Rhodamine 3 B			Rose bengale	
	51. Brown-red ppt.	Slightly darker, on excess brown crystals ppt.	Slightly darker. Ppt.	Ppt.	122. Brown-red ppt.	Ppt.	Brown-red ppt.	Brown-red ppt.	Brown-red flocks.	Conc. sol. light-red flocky ppt.	Conc. sol. flocky ppt.	Ppt. Ppt.	Fuchsine red, slowly becoming turbid.
		2	8 %	{ 109.		139	131.	.133.	134.	135	7140.	145	146.
KED-Continued.	Brillant orange R Scarlet G R, R Xylidhe orange Orange N	Crystal poneenu G. R	Coccluine B. Friku B. 72.	Clayton cloth red	Cloth red G. Cloth red H. Fast red 7 B	Cloth red B Ppt.	Cloth red G, G extra. [131. Brown-red ppt.	Bordeaux B X Brown-red ppt.	Oraclile red A	Cloth scarlet G135, Conc. sol. light-red	Poncean 3 R B. Poncean 3 R B. Poncean 4 Extra Ponceau B. Pow red f. Power of C. Power	Cloth scarlet R	Croceline scarlet 7 II Fuchkine red. slowly Croceline scarlet 8 B. Foreson R. Foreson 6 IR II

KED - Continued.

				Almost no reaction,					
terrate.	No Reaction—Continued. Victoria triby Wool red extra. Oenanthinine	Cochemille red A. Crockine searlet 4 B X. Brilliant searlet. Brilliant ponceau 5 It	Poncenu 4 16 Poncenu brilint 5 16 New coccine Scarlet	88 2	Cotton ponceau 175. Cliniabar scarlet 175. Acid and milling scarlet 240.	Fuchsine 8. Acid magenta Ruthy 8. Acid fuchsine.	Marron S Greinit S Acid cerise	Alizarine V I Alizarine No. I. Alizarine I V. Alizarine No. I B new.	Alzarine P. Alzarine for vlolet. Alzarine unavon.in puste
MBD :- Continuen.	364. Decolorized, and red-	365. Plesh-colored Hocks.	451. Ppt.	38		61.	£ 8	95.	\$H5.
	RED—Continued. Phloatine lodine free Phloatine I A. Decolorized, and red- production of the free 1984. Decolorized, and red-	Grantee iten Cyanoshie Kosino 10 B. Rose bengale Rose bengale Rose bengale Rose bengale	Azo carnine G in paste	No Reaction: Chromotrope 2 R	Brilliant ponceau G Ponceau J. Ponceau B. 4 It Aro Bordeaux	Printalo (109) Fract red B. Bordenux B. B. G. Corneline Corneline Fractor	Chromotrope 10 B. Azo Cochenille. Rast red E.	Acid carmoisine B. Crocelne 3 B X. Frat red D, E B, N S.	American Ago acid ruly 2 B. Bordoux D H, S. Pourpre. Nanhthol red S. 0

			Blue-violet.		Red-violet.	Ppt,		Red-violet ppt.				Turns bluer. Turns bluer.		30. Brownish.	Ppt., water soluble.		Ci Vollous bacours	Tellow Diowin.	62. Light-brown.	
,		_	, 59B.	_		450.		472.		_		: : 48.				. 60.				
ViolET-Continued.	Alizarine S D G, N	Flavo purpurme	Alizarine S X, S X extra	Authra purparine	Alizarine Bordenux B in puste 207 403. Alizarine Bordenux B D in puste 403. Alizarine Bordenux G. G G.	Induline scarlet450.	Rose Magdala (Magdala red) Naphthylamine pink	Naphtbaline red \ 472.	Naphthaline pink Naphaline scarlet	Soudan red.	BLUE:	Azo fuchsine (f 87.	Phones	Orseille substitute 3 V N	Apollo red	Fast red B T 60.	Fast red BBordeaux B. B L. G. B extra	Cerashe Description	Crystal Poncenu 6 R. Azo rosine 677	Coccinine B
	r solution.		Ppt., on boiling red sol.	,	Red-violet ppt., water soluble,		nlcohol sol. red-violet.		Red-violet.	Conc. sol. pyt., sol-		Dirty violet, in conc. sol. dirty-	(violet ppr.	Red-violet ppt.,						
RED.	On addition of NaOH to water solution.	VIOLET:	Azarine S (66. Ppt., on boiling red	5 - 11-5	Cloth red v. (192, Red-violet ppt., water Cloth red f. B.)	Croceine B	Fart Poncentt Tel-violet, Fart Poncentt Tel-	Crootine 3 B	Fast scarlet B	Fast Ponceau 2 B	Bordeaux G116.	Croceine scarlet 7 B cone. Croceine scarlet 8 B cone. Croceine scarlet 8 Croceine cone of the cone.	l'oncean o ft 13	Hessian purple B	Alizarine V 1	Alizarine Ic	Alizarine P	Alizarine marron in paste, 384.	Alizarine W S powder Alizarine WW S worder Alizarine worder W S Alizarine W S Alizarin	Alizarine carmine

RED-Continued.

Yellow - brown ppt., soluble in alcohol or ether with strong fluorescence,	° :300. { Almost colorless, busic ppt.	310. { Blenched almost com-	Bleaches slowly. On heating bleaching with basic ppt.	Fuchsine red, rapidly bleaching on exposure to light or air.		Turns yellower,
Brown—Continued. Neutral red		Fuchsine S. Acid ungenta Ruby Acid furchaine Fuchsine acide Marron S. Greand S.	Rhodamine S	Purpurlue in paste	YELLOW: Soudan G 18. Cerasin uninge 18.	Cocuentitie Scarlet Gr
94 95. Yellow-brown.	97. 18. Red-brown. (94.) Red-brown ppt. sol-	Turns browner,				Mater soluble.
:	Cochemin Pad A. Cochemin Pad	Fast acid Pouceau Powent acid 6. Fast ncid scribe. Brilliant crocefne M Brilliant crocefne buish. Cotton scalet.	Colcol Rearies of Source Colcol Rear Service binish. Paper scribt binish.	Brythrine X. Oreelle red A. Union fast cluret.	Goots service Co. 1535. Cloth scarlet R. 143. Bordeaux B.X. 147. Brilliant croccine 9 B . 156.	Congo Congo

Brown-red ppt. Blue-red. Yellow-red.	Brick-red.	On bolling.	Bluish-red ppt.	Turns darker.	Turns darker. Slightly darker.	Turns turbid and darker,	Ppt. { Gonc. sol. brown red
## 4 9		66.	21	99.	.101	\	127 133 137 138
UED: Orseille substitute V	Woolscarlet R	Azarine R	Rouge d'oxy-amido-diphenne 72. Brika B. 72. Fast red D, E B, N 8.		Wool rete extra. Craatthinine. Roxumine. Fast red B. 101.	First red A First red O First red O	Cloth red III B extra
				yel-	-los inwui	ŧ.	
37. Brown-yellow.	Turns yellower.	Ppt.	Turns yellower.	93. Turns slightly yel-	Red-yellow ppt. sol- uble in large quan- tity of water. Turns yellower.	Brown-yellow p	Orange-yellow, Orange yellew, Orange-red. Orange-red.
55	20.	53. Ppt.	59.	93.	210.	.356	21. 12. 181. 181. 258.
Azo Coccino 2 R. Double scarlet R.	Palatine scarlet	Ponceau 3 R	Paintine red Azo red Azo red A	Azo ruby A, S Azo acli ruby Azo acli ruby Azo fuelisine aclde Prat red C Garmoleiue Brilliant Carmolsine O	Rouge rubis A. Brilliant congo R. 210. Acid rosamine A. 3350.	Violantine G	18. Southan G. 18. 18. Cerasin crange 21. Cochenille scarlet G 21. Isouge M 18. Isouge M 18. Isouge M 18. Itouge M 18. Itouge M 18. Itouge M 18. Itouge M 18. 19.

RED - Continued.

I Light red crystal ppt. on boiling. Pale red ppt. Pale red ppt. Pale red ppt. Propt. Propt. Propt. Propt. Propt.	Led flocks. If barted with little NauOH; excess of NaOH, if excess of NaOH, if the little in ether and behavior, on hearthy notice dimethylar unite smell.	I coud no reaction, on warning suponification with color have ppt. Turns darker with dark-green fluorescence.	960. Deeper blue-red.	Fuchsine red, rapidly decolorizing. Searlet-red ppt. Scarlet-red ppt.
RED—Continued.	Rhodamine B	Hodamine 3 B Anisoline Alcohol soluble cosine Primrose à l'alcool D 11 Methyl cosine		Purpurine in maste
	Deeper red. Deeper red. Ppt. Yeliow-red ppt. Ppt.	Ppt. Ppt. Piocky ppt. Ppt.	oo, stid ppt., soudle in water. Carmine-red ppt. Dueper bine-red. Orange-red. Brick-red ppt.	Orange-red. Binish-red. Reddish crystal ppt. of base.
Continued. Ponceau 3 R B. Ponceau 8 Arta Fast Ponceau B Swared L Searlet B G Imperial searlet Imperial searlet B G Institute B Biebrich searlet Backer	Orecilline B B. 144. Cotton Porceau. 715. Clinnubar searlet. 716. Glycine Corinth. 174. Glycine et and the search of James and properties of B. 177. Benzo purpurine d B. 203. Clonge Districtie & B. 205.		25 - 25 - 25 - 25 - 25 - 25 - 25 - 25 -	

Almost no renetion.			Almost no reaction,	- 213. Almost no reaction 225. 240.	Very slight reaction.	If cold, no reaction.	346. { Little NaOH, if cold, no veaction.	No reaction, if cold.
	38.	. 505.	204.	225. 240.	331.	55.	346.	347.
Nu Rkacrios - funtinued. 126. 1	Diamine fast red F	Benzo purpurlie 4 B	Beuzo purpurine B	Diamine red N () 225, Naphtthylene red () 240,	Cornlline Red cornline Aurine B. Paconine.	Rhodamine G extra	Mydalline B extra Ricidanine O. Ricodanine O. Ricolanine extra B Safranille	Rhodamine 3 B Anisoline Anisoline
Brown-red ppl.								
-101.		2		3	<i>i</i>		.105.	119.
Software Sof	NO HEACTION: Sudan I. Curmine naphtha	Chromotrope 2 II	Scribert Orange R. Scribert G. R. Scarlet R. Scriber R. Scriber R. Scriber G.	Orange A. Poncean 2 R. Poncean G. Poncean G.	Ponceau, R. Brilliant Ponceau († Ponceau J.	Azo Bordeaux Buffalo ruby Azo cochenille	Double acarlet extra S. Double brilliant searlet 3 R. Eenrhate brilliant e. Belliant Poucesu 4 R.	Double scarlet 2 R

RED—Continued.

	To alcohol sol.	
No Reaotton—Continued. Brythrosine D Erythrosine B Erythrosine B Erythrosine B Erythrosine B Frytosine B Iodine cosine B Iodine cosine B Iodine cosine B Evimose soluble Evimose soluble Evimose soluble Evimose bluttre Evimo bluttre Bosine bluttre Bosine J Diunthine B		Rosinduline 2 B bluish.
	Isosine B J F F Isosine B N Netyyl cosine Estimate B Solution searlet B Estrate J Solution searlet B Solution searlet B Estrate J Solution searlet B Searlet B Searlet B Solution Searlet B Searlet	Jaune d'Orient

LACK.

Brack: Phenylene black. Authracije hjark 3.	<u> </u>	GREEN: Victoria black B	reen,
0	Blue-black. Greenish-black.	Blue black B	
Diamine blue-black E. Noirblen	-228. Blue-black.	Naphthol black B	
Diamine E.		Diamond black	sh.
Diamine black B O	Bine-black,	Naphthol blue-black	
Fast black '438. Almost black.	Almost black.	285. (D)	Dirty yellow - green, on heating carnois-
BROWN:	Dirty red-brown.		lution of SO ₂ .
Alizarine black S402.	With decomposition.	Fast black B356. Slightly	Slightly soluble with dirty-green color.
Y		Fast black B S	Paste, if dry, dirty-
Anillne black		Nigramine501. Gray.green.	reen.
Aniline black in paste		New gray	
		New methylene gray G B. Nigrishne. Malta gray G B. Gris direct J. Ben pouldre.	· · ·
Violet black 173. Diamino black R O. 187. Direct gray R. 187. Direct gray B. 224.	Bluish. Bluish.	Gris direct 4 R. R on pâte. Gris spécial R en pâte. Alsace gray	
Diamine blue-black E Spiraline E Sinck-blue friamine E		383	Dirty yellow - green,
Diamine black B O	Black-blue,	Naphthazarine S	ine-red with evo- lution of SO ₂ . irty brown-red.

BLACK -- Continued.

BLACK			VIOLETT: Naphthol black 6 H	Green-blue, then vio-	,0
On addition of H ₂ O to H.SO, solution.	SO	solution.	Naphthol black B 187. Blue ppt., then red-livilliant black B. violet ppt.	Blue ppt., then red-	
Activity Place Direct. (152). Greenish-black ppt. Authorette black II. (154).	132.	Greenish-black ppt.	Digmond black	Ppt. Ppt.	
Naphthylamine black D1531, Green, then black	1531.	Green, then black	First black. First blue-black in paste	Hack-violet.	
Menzo gray	355.	Ppt. [Brownleh sol, and [black ppt.	HLUK: Jot black R. 150. Greenish-blue prt. Naphthol black 6 B. 154. Green blue, then	Greenish-blue put. Green - blue, then violet met.	
Frak black Frak black in paste 4.78. Violet-black.	<u> </u>	Violet-black.	Azo bluck B 186. Blue sol. and bluo ppt.	Blue sol. and blue ppt.	
New gray Methylene gray O. N D. N F New methylene gray G. B.			Augustrol ones, 5. Frilliant black B. Naphthol blue-linek. Naphthol 12 B. Naphthol 12 B.	Die ppe, caea rec- violet ppt. Ppt.	
Matta gray. Orla direct J. B en pondre. Orla direct J R, R en pûte.	500g	Reddish-gray.	Diamlno black R O. 187. Direct gray R. 117. Direct gray H. 221.	Bed-blue ppt. Groy-blue ppt. Gray-blue ppt.	
Abaco gray			Pilamine blue-black E	Ppt.	
Over State S	1330.	Ppt., decomposes on bolling.	Dhambe black H O 229t. Red-blue ppt. Noir Diamibe H Mark Market Mar	Red-blue ppt.	
Alizarine black 8 in paste	376.	Brownish sol, and black ppt.	Gnk	On large dilution blue-est.	
	÷ 0.1.	Light brown sol., on cooling brown ppt.	Jot black R	Bluish-green ppt. Dark-green ppt. of sulphate.	

Ppt. Ppt.	Buistt-black ppt. Ppt. Ppt.	Pyt. Dark-blue ppt. Dark-blue ppt. Ppt.	Ppt. Blue-black ppt.	Gray-blue.	Ppt. (soluble in alkn-lies) with evolution of H ₂ S.	Bordeaux red ppt.
Violkt—Continued, Dlamoud black, Violet black	BLUE: Jet black IL Jet black IE Blaish-black ppt. Hue black B Azo black O 155. Ppt. Naphtol blue black Noir uspitch of 2 B	Direct gray B. 0. 187,	438.	New gray Neth plene gray O, N D, N F New methylene gray G, B. Nigrisine Malta gray Gris direct J, B en pondre Gris direct J, B en pite. Gris direct R en pite. Alouve gray	OREEN: Noir vidut Noir vidut (494.) (494.) (494.) (494.) (494.)	Red); Victoria black H
Green, on large dllu-	Reddish-gray.	Solution.		l'pt. Bluc-black ppt. Brown-red.	Red-vlolet ppt.	Deep violet ppt. Ited-violet ppt.
RED: Victoria black B. Victoria black G, 5 G. Victoria black G, 5 G. Victoria black G, 6 G. Victoria black G.	Now gray	Alsace gray	BLACK : 150, Joe black R. Napitthylamine black D. 151, 153, Bears gray. 211, Bears gray.	Alizarine black S. Fast black Frat black in paste Brown: Alizarine black S in paste. Alizarine black S W in paste.	Alizarine black P	Anthraele black II. Naphthol black II. 152. Deep violet ppt. Naphthol black II. 157. ted-violet ppt. 157. ted-violet ppt.

BLACK - Continued.

		23. Illnek-vloiet ppt. 21. Ppt. 31. Historiak	ib. Ppt.	7. Ppt., water soluble.	ii. ir.	I. Strong dirty-green.	(Or in rodn sol.), if cold incoluble, on heating turning to violatived,		e
BLACK - Continued.	Viouxy	Alizaribe black S	Hine-black B. Azo black O.	Naphthol black B	Diamond black	Chuic	Uku: Fast black B	No IRA-centric: Naphtdynanine black D. 153, Naphtdynanine black D. 163, Noir naphtdol 12 B. 164, Diamine blace B. 165, Noiriber B. 165, Dannine B. 165,	Diamine black B O
PLACK	385. Brown-red.			. solution.	Muc-black, Violet-black ppt.	Grny - black put, of the base, sciuble in benzole or ether,	with cherry-red color. On addition of water or needle neid blue-green sol.	Ppt. Dark blue-violet.	Dark blue-violet. Beep-violet ppt.
	Ukro - Confibured. Alfardine black S in paste Alfardine black S W in paste Alfardine black B W in paste Alfardine black B W	No Reaction : Naphthol black 6 H	BLACK.	On addition of NaOH to water solution.	Heack: Napithol black 6 B.: 154. Blue-black. Frat black: Frat blueblack in paste. 448. Violet-black ppt.	New grny Mathylene grny O, N B, N F New Mathylene grny, Q, 11 Nigrildne	Martin gray. Martin gray. Gris direct 3, II en pandre. Gris direct 4 It, II en palte. A less greint, II en pâte.	VIOLET : Wood black Nord pour laine Victoria black II. Victoria black II.	Jet black R. Dark blue-violet. Phenylene black B. Beep-violet ppt.

PART X. — ALPHABETICAL LIST OF DYES, GIVING AGENT AND DYEING METHOD.

ABBREVIATIONS USED TO INDICATE THE DYE MAN-UFACTURERS AND THEIR AGENTS.

- (A A C) American Color & Chemical Co., Albany, N. Y. F. E. Atteaux & Co., 174-176 Purchase St., Boston, Mass.
- (A) Action-Gesellschaft für Anilinfabrikation, Berlin, Germany. New York and Boston Dyewood Company, 55 Beekman St., New York.
 - (AC) Anchor Color Mfg. Co., 462 Cherry St., New York.
- (APM) Alex. P. Mende, 536-540 West Fourteenth St., New York.
- (A M) Actiengesellschaft f\u00fcr Chemische Industrie in Mannheim, Germany.
 - (At) F. E. Atteaux & Co., 174-176 Purchase St., Boston, Mass.
- (B) Badische Anilin und Sodafabrik, Ludwigshafen am Rhein, Germany. Pickhardt & Kuttroff, 128 Duane St., New York.
- (BA Co.) The British Alizarine Company, Ltd., Silvertown, Vietoria Docks, London, England. Beach & Co., Hartford, Ct.
 - (Bai) Bairstow & Co., 211 Pearl St., New York.
- (BCF) Baseler Chemische Fabrik, Bindschedler, Basle, Switzerland.
 - (Beh) Beach & Co., Hartford, Ct.
 - (Bd) J. A. & W. Bird & Co., 43 Cedar St., New York.
 - (BE) C. vom Bauer in Elberfeld, Germany.
- (BK) Leipziger Anilinfabrik Beyer and Kegel, Leipzig, Lindenau u. Fürstenberg-on-Oder, Germany.
 - (BL) Bosson & Lane, 36 Central Wharf, Boston, Mass.
 - (Bt) F. Brett & Co., 194 Fulton St., New York.

- (Br S) Brooke, Simpson & Spiller, Ltd., Atlas Works, Hackney Wick, London, England. Beach & Co., Hartford, Ct.
 - (Bs) C. Bischoff & Co., 87-89 Park Place, New York
- (By) Farbenfabriken vormals Fried. Bayer & Co., Elberfeld, Germany. Farbenfabriken, 40 Stone St., New York.
- (C) Leopold Cassella & Co., Frankfort-on-Main, Germany. Wm. J. Matheson & Co., Ltd., 182-184 Front St., New York.
- (CFG) Chemische Fabrik Griesheim, Frankfort-on-Main. Germany.
- (Cl Co) The Clayton Aniline Company, Ltd., Clayton, Manchester, England. J. A. & W. Bird & Co., 43 Cedar St., New York.
 - (CJ) Carl Jäger in Barmen, Barmen, Germany.
 - (CR) Claus & Reé, Droylsden, Manchester, England.
- (C V) Colne Vale Chemical Co., Milnsbridge, near Huddersfield, England.
- (Cz) John Casthelaz, Bruère & Co., Belbeuf, near Rouen, France.
- (D) Farbenfabrik Dahl & Co., Barmen near Elberfeld, Germany. C. Bischoff & Co., 87-89 Park Place, New York.
- (DH) L. Durand, Huguenin & Co., Basel, St. Fons and Hüningen, Germany. A. Klipstein Company, 122 Pearl St., New York.
- (DW) L. Destree, A. Wiescher & Co., Haeren bei Brüssel, Belgium.
 - (EH) E. de Haën in List vor Hannover, Germany.
- (E F W) Elton Fold Works, Bury, Lancashire, England. Beach & Co., Hartford, Ct.
- (F) Farbwerk Friedrichsfeld, Dr. Paul Remy in Mannheim, Germany.
- (Fi) Alfred Fischesser & Co., Lutterbach bei Mulhausen,
- (FTM) Fabriques de Produit Chimiques de Thann et de Mulhouse, Alsace.
- (G) Joh. Rud. Geigy & Co., Basel, Switzerland. John J. Keller & Co., 104-106 Murray St., New York.
- (Gau) Gauhe & Co., Alizarinfabrik in Eitorf a. d. Sieg, Germany.
- (Gb) Anilinfarbenwerk vorm. A. Gerber & Co., Basel, Switz-
 - (Gei.) Geisenheimer & Co., 189 Front St., New York.

- (Gr.) Rob. Graesser, Chemical Works, near Ruabon, North Wales.
- (H) Read Holliday & Sons, Ltd., Huddersfield, England. Read Holliday & Sons, 100 William St., New York.
 - (H M) The Heller & Merz Co., Newark, N. J.
- (H R W). Hudson River Aniline Color Works, Albany, N. Y. Farbenfabriken of Elberfeld Co., 40 Stone St., New York.
- (H S) The Hanna-Schœllkopf Co., 135 North Water St., Philadelphia.
- (I) Gesellschaft für Chemische Industrie, Basel, Switzerland. A. Klipstein & Co., 122 Pearl St., New York.
 - (Jb) J. B. Ibels, Chausée de Mons, Brüssels, Belgium.
 - (J H) J. Hauff in Feuerbach, near Stuttgart, Germany.
 - (Jy) O. S. Janney & Co., 8-10 Letitia St., Philadelphia.
- (K) Kalle & Co., Biebrich-on-Rhine, Germany. Kalle & Co., 77 John St., New York.
- (KB) Küchler & Buff, Crefeld, Germany. F. Bredt & Co., 194 Fulton St., New York.
 - (Kchl) Victor Koechl & Co., 122 Hudson St., New York.
 - (Kell.) John J. Keller & Co., 104-6 Murray St., New York.
 - (Ki) Kinzlberger & Co., Prague, Hungary.
 - (Klp) A. Klipstein & Co., 122 Pearl St., New York.
- (L) Farbwerk Mühlheim vorm. A. Leonhardt & Co., Mühlheim-on-Main, Germany. Victor Koechl & Co., 122 Hudson St., New York.
- (L D) Lepetit, Dollfus e Gansser, Susa, Milan, Italy. J. A. & W. Bird & Co., 43 Cedar St., New York.
- (Lev) Levinstein Limited, Slackle and Crumpsall, Manchester, England.
 - (LM) Leeds Mfg. Co., Brooklyn, N. Y.
 - (Lo) Charles Lowe & Co., Stockport, England.
 - (LP) Lucien Picard & Co., St. Fons, Lyons, France.
- (L Sch) Lembach & Schleicher, Biebrich-on-Rhine, Germany.
- (M) Farbwerke vorm. Meister Lucius & Brüning, Höchston-Main, Germany. Victor Koechl & Co., 122 Hudson St., New York.
- (Math) Wm. J. Matheson & Co., Ltd., 182-184 Front St., New York.
- (M Ly) Manufacture Lyonnaise des Matières Colorantes, La Mouche, near Lyons, France. Wm. J. Matheson & Co., Ltd., 182-184 Front St., New York.

(Mo) Gillard P. Monnet & Cartier, St. Fons, Lyons, France.

(N) Carl Neuhaus, Elberfeld, Germany. A. Klipstein & Co., 122 Pearl St., New York.

(NY & B) New York and Boston Dyewood Co., 55 Beekman St., New York.

- (N I) Farbwerke Griesheim, W. Noetzel, Istel & Co., Griesheim-on-Main, Germany.
- (O) K. Oehler, Anilin u. Anilinfarbenfabrik, Offenbachon-Main, Germany. Geisenheimer & Co., 189 Front St., New York.
- (P) Société Anonyme des Matières Colorantes et Produits Chimiques de St. Denis, France. Sykes & Street, 85 Water St., New York.
- (P) St. Denis Dyestuff & Chemical Co. (Λ. Poirrier), Paris, France. Sykes & Street, 85 Water St., New York.
 - (PC) Theodor Peters, Chemnitz, Germany.
 - (PK) Pickhardt & Kuttroff, 128 Duane St., New York.
 - (P L) Pick, Lange & Co., Amersfoort, Holland.
- $({\rm P}\,{\rm S})\,$ Ferd. Petersen & Co., Schweizerhalle, Basel, Switzerland.
 - (R) Joh. Conr. Reihlen, Friedensau, Neuhofen, Germany.
- (R D) Roberts, Dale & Co., Manchester and Warrington, England.
- (R E) Dr. Remy & Co., Weissenthurm-on-Rhine, Germany. Fr. Schroeder, 100 William St., New York.
- (RF) J. Ruch et Fils, Pantin (Seine), maison à Paris, France.
- (Rh) Société Chimique des Usines du Rhone, France. Beach & Co., Hartford, Ct.
- (S) Chemische Fabrik vorm. Sandoz & Co., Basel, Switzerland.
 - (SB) A. Sevoz & Boasson, Lyon-Vaise, France.
 - (S M) The Stamford Mfg. Co., 133-137 Front St., New York.
- (Sch) The Schoellkopf Aniline and Chemical Co., Buffalo, N. Y., 100 William St., New York.
 - (SS) Sykes & Street, 85 Water St., New York.
 - (S W) The Stone & Ware Co., 157 Maiden Lane, New York.
 - (T) W. G. Thompson & Co., Middleton.
- (tM) Chemische Fabriken vorm. Weiler-ter Meer, Uerdingen, Germany.
 - (V) Verein Chemischer Fabriken in Mannheim, Germany.

The Roessler & Hasslacher Chemical Co., 100 William St., New York.

- (W) Williams Brothers, Hounslow, Middlesex, England.
- (W C B) W. C. Bearnes & Co., Ltd., Hackney Wick, London, England. Wm. J. Matheson & Co., Ltd., 182-184 Front St., New York.
 - (Z) Friedrich K. Zimmer, Mannheim, Germany.

DYEING METHODS.

The dyeing formulæ used throughout are referred to by number. This is done to save space, and leaves the dyeing formulæ all in one place.

For the reason that the records are meagre and faulty, this list is not offered except E. & O. E. The authorities vary and the processes themselves alter and improve. We will thank any one for the courtesy of corrections and additions.

WOOL.

- 1. Neutral bath, as magenta, methyl violet, etc., begin at 40° Centigrade, and after bringing to a boil very slowly, add afterward 1 per cent. acetic acid, to brighten the color.
- 2. Neutral bath, as diamin fast red, sulfonazurine. Dyed with 15 per cent. Glauber salts, with the least possible amount of water.
- 2a. Colors dyed with 3 per cent. acetic acid, viz., lanacyl blue.
- 3. Colors dyed in an acid bath, as acid fuchsin, ponceau, fast red, with 15 per cent. of Glauber salts. Commence 70° Centigrade, add slowly 5 per cent. sulphuric acid, previously diluted with water.
- 3a. Ten per cent. Glauber salts; enter hot; add 2 per cent. bisulphate; after fifteen minutes, 8 per cent. of bisulphate.
- 3b. Dye with Glauber salts, sulphuric acid and chrome afterward.
- 4. Chrome mordant. Boil an hour and a half with 3 per cent. bichrome and $2\frac{1}{2}$ per cent. tartar. Tartar is often replaced by $1\frac{1}{2}$ per cent. of sulphuric acid, or sodium bisulphate, or oxalic acid. Acetic acid may aid in exhausting the bath.

- 4a. Mordant with 11/2 per cent bichrome, 3 per cent. lactic acid of 50° Beaumé, and 1 per cent. sulphuric acid.
- 5. Chrome mordant with fluorchrome. Boil an hour and a half in a wooden vat with 4 per cent. fluorchrome and 1 per cent. oxalic acid.
- 6. Mordant dyes. Dissolve the dyes in water containing 10 per cent. of acetic acid. Enter the goods at 50° Centigrade, carry to a boil, after half an hour add slowly 2 per cent. of bichrome and 2 per cent. of tartar in solution; boil again three-quarters of an hour.

6a. Six per cent. copperas, 3 per cent. cream of tartar, 1 per cent. oxalic acid.

6b. For colors that dye on an alum mordant. Mordant the wool with 12 per cent, alum and 7 per cent, cream of tartar.

SILK.

Process 7. Dyes which are put on in a neutral bath slightly acidified with acetic acid, as magenta, methyl violet, etc .-For every 5 kilos of silk used take 120 liters of water, 20 liters of soap solution with acetic acid up to a slightly acid reaction. Enter at 85° Centigrade; wash after dueing and vivify with about 20° c.c. of acetic acid in 120 liters of water. Dry without washing. The soap solution is the soap solution used in boiling out the silk, to which is added one-third of its weight of a pure Kern soap. German dyers added to this soap solution 20 kilograms of Marseilles soap, 200 grams of gelatine, 10 grams of common salt in every 100 liters of water. French dyers use starch paste instead of gelatine.

Process 8. Dves which are used in an acid bath as fast red, ponceau, naphthol yellow S. etc.-For every 5 kilos of the dve use 120 liters of water, and 30 liters of the soap solution. Wash and vivify with about 10 c.c. of sulphuric acid in 120 liters of water.

Process 9. Chrome Mordant.-In chrome chloride or 20° Beaumé leave the silk for twelve hours at 40°. Wash well and dry out in the centrifugals. Pass into a water-glass solution of 1° Beaumé and turn five times. Dye in the soap solution, to which acetic acid has been added. Wash, boil in a soap solution containing 5 gm. of Kern soap, wash, and vivify with lukewarm water with acetic acid.

Process 10. Alum Mordant.—Take 100 gm. of iron-free alum, 40 gm. of hyposulphite of soda (Na₂SO₂) in a liter of water, one hour at 40° Centigrade, one hour at 60°, and one hour at 80°. Wash and dye in a soap solution which has been slightly acidified with acetic acid in the same way as in Process 9a.

Process 10b.—Sixty gm. of iron-free alum, 6 gm. of soda crystals, 1 liter of water; precipitate which is formed first is redissolved, turn four or five times, and give twelve hours. Put on the centrifugals, then in water-glass solution of 1° Beaumé, wash carefully, and dye. The dye-bath should receive 20 liters of soap solution for every 100 liters of water, and the bath should be slightly acidified with acetic acid, after which it is dyed as in process 9a.

COTTON AND LINEN.

Process 11, Adapted for magenta, methylene blue, etc.—Mordant in 5 per cent. tannin, calculated on the weight of the goods, using very little water, enter at 50° or 60° Centigrade, let it stand for six hours; place in the centrifugals, and then enter into a bath of 20 per cent. tartar emetic at 30° Centigrade for fifteen or twenty minutes; lift, and wash thoroughly.

Process 12. Direct dyeing substantive dyes, as chrysamine, chrysophenine, diamine blacks, etc.—Dye with 3 per cent. of soap, 20 per cent. of common salt or Glauber salts. Wash and dry. Hard water must absolutely be avoided. Phosphate of soda is used in the dye-bath; also soda, or potash, or waterglass.

Process 12a.—Forty per cent. of salt and 2 to 5 per cent. of soda-ash.

Process 12b.—Forty per cent. of Glauber salts and 2 to 5 per cent. of soda ash.

Process 12c.—Glauber salts

Process 12d.—Glauber salts and soap.

Process 12e.—Use soap alone.

Process 12f.—Use salt only.

Process 12g.—Alum and Glauber salts.

Process 13. Chrome Mordant.-Allow to stand twelve hours in the chrome mordant of the Höchst Dye Works, then wring, and place for half an hour in a solution of 30 gm. of soda-ash per liter at 60° Centigrade. Wash well.

Process 14.—Turkey red oil, sumac, alum, chalk.

Process 15.-Diazotize and couple with

- a. Alpha-naphthol.
- b. Beta-naphthol.
- c. M-phenylene-diamine.
- d. Toluylene diamine hydrochloride.
- e. Diamine hydrochloride.
- f. Diazotized benzidin.
- g. Diazotized paranitraniline.
- h. Dve with 15 to 20 per cent. Glauber or common salt, and then pass into diazotized azophor red, etc.
- i. Resorcin.
- j. Amido-naphthol ether.
- k. Nitrazol.

Process 16.—Ten per cent. sulphoricinate of soda, with 0.2 per cent. stannate of soda; dry and enter into acetate of alum of 5° Beaumé. Dry at 40° to 50° Centigrade. Dye with 1 per cent. acetic acid.

Process 17.-For dyes that dye on aluminum mordanted cotton or linen, prepare the mordanting solution the following way: 1 liter of water, 200 gm. of normal aluminum sulphate, 31.82 gm. of sodium carbonate (Na₂CO₂); make the solution to stand at 10° Twaddle (sp. gr. 1.05). Impregnate the cotton with this solution; remove excess of liquid by squeezing or wringing; dry at a low temperature, and pass into cold H.O containing 50 gm, of strong commercial ammonia liquor (0.88 sp. gr.) per liter. Work in this solution for ten minutes and wash well.

If soap be the fixing agent employed, use a solution containing 10 gm. per liter.

If arsenate or phosphate of soda, use from 5 to 10 gm. per liter.

If silicate of soda at 100° Twaddle, use 5 to 10 gm. per liter.

ALPHABETICAL LIST OF DYES.

Name of Dye.	Agent.	Method.
Acetine Blue	B	
Acetinduline R solution		
Acid Alizarine Blue B B and G R		3
Acid Alizarine Green B and G	M	3
Acid Black B		3
Acid Black B, No. 4	A C	3
Acid Black No. 77		3
Acid Black No. 5,534	Н	3
Acid Black No. 5,535	Н	3
Acid Black S		
Acid Blue A A	G	3
Acid Blue 6 G		3
Acid Blue 100		3
Acid Brown	D	3
Acid Brown G	A	
Acid Brown R	A	
Acid Carmoisine B	В К	3
Acid Cerise	, P, P S	
Acid Fuchsine	D. F, L	3-8
Acid GreenBy; D	H, F, O	3-8
Acid Green		
Acid GreenBy, I	F, O, tM	
Acid Green bluish		3—8
Acid Green D		
Acid Green extra conc	C	3-8
Acid Green M		3-8
Acid Green 3 B	\dots By	3
Acid Green B, 2 B, 3 B, 4 B		3
Acid Green J		3
Acid Green J E E E		3
Acid Grenadine		3
Acid Grenadine B		3
Acid Indigo Blue	Sch	3
Acid MagentaB,	PS. SS	
Acid and Milling Scarlet	\dots Br S	3
Acid Methyl Violet S 7 B	P K	3
Acid Naphthol Orange		3
Acid Orange	G	3-8
Acid Ponceau		
Acid Rosamine A	M	3-8
Acid Rubin (see Fuchsine S)		3-8
Acid Rubin S B	F K	3
Acid Ruby		2
Acid Violet 2 B	B	
Acid Violet 2 B		3
Acid Violet 3 B N	В	

Acid Violet 3 B extra By 3 Acid Violet 4 B N. By 3 Acid Violet 4 B N. By 3 Acid Violet 5 B. S S 3 Acid Violet 5 B F. M 3 Acid Violet 6 B. G 3 Acid Violet 6 B. By 3 Acid Violet 6 B. A 3 Acid Violet 7 B. L 3 Acid Violet 7 B. B. J Acid Violet 2 R extra. By 3 Acid Violet 2 R extra. A B Acid Yellow B. A A 3 Acid Yellow B. A A 3 <t< th=""><th>Name of Dye.</th><th>Agent.</th><th>Method.</th></t<>	Name of Dye.	Agent.	Method.
Acid Violet 4 B extra By 3 Acid Violet 4 B N B, I 3 Acid Violet 4 B N By 3 Acid Violet 5 B S S 3 Acid Violet 6 B G 3 Acid Violet 6 B By 3 Acid Violet 6 B A 3 Acid Violet 6 B N B, I 3 Acid Violet 7 B L 3 Acid Violet N M 3 Acid Violet 2 R extra By 3 Acid Violet 3 R extra By 3 Acid Violet 4 R B A Acid Violet 4 R B A Acid Violet 4 R B A Acid Yellow A, F, P, P, S, R, F Acid Yellow B A A Acid Yellow B A A Acid Yellow B B A Acid Yellow S (see Naphthol Yellow S) A Acid Yellow R B A Acridine Orange G B B Acridine Orange			
Acid Violet 4 B N. B, I 3 Acid Violet 5 B. S S 3 Acid Violet 5 B F. M 3 Acid Violet 6 B. By 3 Acid Violet 6 B. By 3 Acid Violet 6 B. By 3 Acid Violet 6 B. B, I 3-8 Acid Violet 7 B. L 3 Acid Violet 7 B. B, J 3-8 Acid Violet 7 B. B, J 3-8 Acid Violet 8 Extra. By 3 Acid Violet 1 R Extra. By 3 Acid Violet 4 R. By 3 Acid Violet 4 R. By 3 Acid Violet 4 R. By 3 Acid Yellow D. A 3 Acid Yellow B. A 3 Acid Yellow G. D 3 Acid Yellow R. D 3 Acid Yellow, 8,522 H 3 Acridine Orange R. L, S.B. 11-7 Acridine Orange R. L, S.B. 11-7 Acridine Orange R. L, S.B. 1			
Acid Violet 4 B X By 3 Acid Violet 5 B F M 3 Acid Violet 6 B G 3 Acid Violet 6 B By 3 Acid Violet 6 B A 3 Acid Violet 7 B L 3 Acid Violet 7 B B J Acid Violet 7 B B J Acid Violet 2 R extra By 3 Acid Violet 3 R extra By 3 Acid Violet 3 R extra By 3 Acid Violet 4 R B A Acid Violet 4 R B A Acid Yellow A F, P, P S, R F Acid Yellow B A 3 Acid Yellow G A 3 Acid Yellow R S D 3 Acid Yellow S (see Naphthol Yellow S) A Acid Yellow S, S22 H 3 Acridine Orange G Bs 11 Acridine Orange R O L L Acridine Orange R O L S Acridine O			
Acid Violet 5 B F. M 3 Acid Violet 6 B. G 3 Acid Violet 6 B. By 3 Acid Violet 6 B. A 3 Acid Violet 6 B. A 3 Acid Violet 7 B. L 3 Acid Violet 7 B. B, J 3-8 Acid Violet 7 B. B, J 3-8 Acid Violet 8 R. By 3 Acid Violet 1 R. By 3 Acid Violet 3 R extra. By 3 Acid Violet 4 R. B 3 Acid Violet 7 B. A 3 Acid Violet 8 R. B 3 Acid Violet 9 R. A 4 F, P, P.S. R.F. Acid Violet 4 R. B A 3 Acid Yellow S. C B 1 Acid Yellow R.			
Acid Violet 5 B F. M 3 Acid Violet 6 B. By 3 Acid Violet 6 B. A 3 Acid Violet 6 B. A 3 Acid Violet 6 B. A 3 Acid Violet 7 B. L 3 Acid Violet 7 B. By 3 Acid Violet 8 Extra By 3 Acid Violet 1 R extra By 3 Acid Violet 3 R extra By 3 Acid Violet 4 R B B Acid Violet 4 R B B Acid Violet 4 R B B Acid Vellow A F, P, P S, R F Acid Vellow B A 3 Acid Yellow B A 3 Acid Yellow B A 3 Acid Yellow R B D 3 Acridine Orange R E B 11-7 Acridine Orange R E E<			
Acid Violet 6 B.			
Acid Violet 6 B By 3 Acid Violet 6 B N B. I 3—8 Acid Violet 7 B L 3 Acid Violet 7 B B. J 3—8 Acid Violet 7 B B. J 3—8 Acid Violet 8 Extra By 3 Acid Violet 2 R extra By 3 Acid Violet 4 R S By 3 Acid Violet 4 R S M 3 Acid Yellow A, F, P, P S, R F Acid Yellow D A 3 Acid Yellow R S D 3 Acid Yellow B S B 11—7 Acridine Orange R Extra L			
Acid Violet 6 B. A 3 Acid Violet 7 B L 3 Acid Violet 7 B B. J 3-8 Acid Violet 7 B B. J 3-8 Acid Violet 8 B. By 3 Acid Violet 2 R extra By 3 Acid Violet 3 R extra By 3 Acid Violet 4 R B B Acid Violet 4 R B A Acid Vellow G M 3 Acid Yellow D A A, F, P, P S, R F Acid Yellow G D 3 Acid Yellow S (see Naphthol Yellow S) D 3 Acid Yellow S,822 H 3 Acridine Orange G Bs 11 Acridine Orange R extra L S B 11 Acridine Gange R extra L S B 11 Acridine Scarlet R, 2 R, 3 R L L Acridine Scarlet R, 2 R, 3 R L L Acridine Scarlet R, 2 R, 3 R L L Acridine Scarlet R, 2 R, 3 R L L Acridine Scarlet R, 2 R, 3 R L L			
Acid Violet 6 B N. B. I 3—8 Acid Violet 7 B. B. J 3—8 Acid Violet N. M 3 Acid Violet R extra. By 3 Acid Violet 2 R extra. By 3 Acid Violet 3 R extra. By 3 Acid Violet 4 R. B Acid Violet 4 R. B Acid Violet 4 R. M 3 Acid Yellow G. M 3 Acid Yellow G. D A 3 Acid Yellow G. D 3 Acid Yellow S (see Naphthol Yellow S) D 3 Acid Yellow S (see Naphthol Yellow S) D 3 Acridine Orange L, S B 11—7 Acridine Orange G. Bs 11 Acridine Orange R extra L, S B 11 Acridine Red B, B B, 3 B. L, S B 11 Acridine Scarlet R, 2 R, 3 R. L L Acthyl Blue B F. M M Acthyl Green K B 3—8—11 Acthyl Green K B 3—8—11 Acthyl Green K B 3—8—11			
Acid Violet 7 B L 3 Acid Violet 7 B B J 3-8 Acid Violet N M 3 Acid Violet 2 R extra By 3 Acid Violet 3 R extra By 3 Acid Violet 4 R B B Acid Violet 4 R S M 3 Acid Yellow A F, P, P S, R F Acid Yellow B A 3 Acid Yellow R S D 3 Acid Yellow S (see Naphthol Yellow S) A Acid Yellow S (see Naphthol Yellow S) B 11-7 Acridine Orange G Bs 11 Acridine Orange R extra L, S B 11-7 Acridine Orange R extra L, S B 11 Acridine Scarlet R, 2 R, 3 R L L Acridine Ped B, B, 3 B L, S B 7-11 Acthylene Blue O 0 Acthylene Blue O 0 Acthylene Blue N 3-8-11 Acthyl Green K B 3-8-11 Acthyl Green K B 3-8-11 Acthyl Green <			
Acid Violet 7 B. B. J 3—8 Acid Violet R extra. By 3 Acid Violet 2 R extra. By 3 Acid Violet 3 R extra. By 3 Acid Violet 4 R. B B Acid Vellow R. M 3 Acid Yellow D. A A Acid Yellow G. D A Acid Yellow S (see Naphthol Yellow S) D 3 Acid Yellow S (see Naphthol Yellow S) D 3 Acridine Orange G. Bs 11 Acridine Orange G. Bs 11 Acridine Orange R extra L, S B 11 Acridine Scarlet R, 2 R, 3 R. L L Acridine Scarlet R, 2 R, 3 R. L L Acthyl Blue B F. M M Acthyl Green K B 3—8—11 Acthyl Green A 3—8—11 A Acthyl Green A 3—8—11 A Alcohol Blue S F C.			
Acid Violet N M 3 Acid Violet 2 R extra By 3 Acid Violet 2 R extra By 3 Acid Violet 4 R B B Acid Violet 4 R S M 3 Acid Violet 4 R S M 3 Acid Yellow A, F, P, P S, R F A Acid Yellow D A 3 Acid Yellow R S D 3 Acid Yellow R S L S B 11 Acridine Orange R extra L S B 11 Acridine Orange R extra L S B 11 Acridine Orange R extra L S B 11 Actycline Scarlet R, 2 R, 3 R L L S B			
Acid Violet R extra. By 3 Acid Violet 2 R extra. By 3 Acid Violet 4 R B 3 Acid Violet 4 R S M 3 Acid Yellow D A, F, P, P S, R F A Acid Yellow B A 3 Acid Yellow G D 3 Acid Yellow S (see Naphthol Yellow S) D 3 Acid Yellow, 8,822 H 3 Acridine Orange G Bs 11 Acridine Orange N O L A Acridine Orange R extra L, S B 11 Acridine Red B, B B, 3 B L, S B 11 Acridine Scarlet R, 2 R, 3 R L L Acridine Yellow L, S B 7-11 Aethyl Blue B F M 1 Acthyl Green K B 3-8-11 Acthyl Green K C K Alcohol Blue S C			
Acid Violet 2 R extra. By 3 Acid Violet 4 R B B Acid Violet 4 R S M 3 Acid Yellow A, F, P, P S, R F A Acid Yellow D A 3 Acid Yellow R S D 3 Acid Yellow R S D 3 Acid Yellow S (see Naphthol Yellow S) D 3 Acid Yellow R S H 3 Acid Yellow R S B 11—7 Acid Yellow R S B 11—7 Acid Yellow R S B 11—7 Acid Yellow R S B 11 Acid Yellow R S B 11 Acid Yellow R S B 11 Acid Yellow R S B 1 Acridine Orange R Sextra B 1 Acridine Orange R extra L S B 11 Acridine Orange R extra L S B 1-11 Acridine Scarlet R, 2 R, 3 R L L S B 7—11 Acthyl Blue B F M A 3—8—11 Acthyl Green K B 3—8—11 <td></td> <td></td> <td></td>			
Acid Violet 3 R extra. By 3 Acid Violet 4 R S. M 3 Acid Vellow . A, F, P, P S, R F Acid Yellow D A 3 Acid Yellow R S. D 3 Acid Yellow S (see Naphthol Yellow S) . . Acid Yellow S (see Naphthol Yellow S) . . Acid Yellow S (see Naphthol Yellow S) . . Acid Yellow R S. D 3 Acridine Orange L S B 11—7 Acridine Orange G Bs 11 Acridine Orange N O L L Acridine Red B, B B, 3 B L S B 11 Acridine Red B, B B, 3 B L S B 11 Acridine Scarlet R, 2 R, 3 R L L Acridine Yellow L, S B 7—11 Aethyl Blue B F M 3—8—11 Aethyl Green K B 3—8—11 Aethyl Green K B 3—8—11 Aethyl Green A B, I 3—8 Alcohol Blue B, Br S, By, C R, D, K L, L, R D R Alcohol Blue B, Br S, By, C R, D, K L, L, R D			3
Acid Violet 4 R			
Acid Violet 4 R S.			3
Acid Yellow D A 3 Acid Yellow G Acid Yellow R S Acid Yellow S (see Naphthol Yellow S) Acid Yellow, 8,822 Acridine Orange G Acridine Orange R extra <			
Acid Yellow B A 3 Acid Yellow R S D 3 Acid Yellow S (see Naphthol Yellow S) B 1 Acid Yellow, 8,822 H 3 Acridine Orange L, S B 11—7 Acridine Orange G Bs 1 Acridine Orange R extra L, S B 11 Acridine Red B, B B, 3 B L, S B 11 Acridine Scarlet R, 2 R, 3 R L L Acridine Yellow L, S B 7—11 Acthyl Blue B F M M Acthyl Green K B 3—8—11 Acthyl Green K B 3—8—11 Acthyl Green A S—8—11 A Acthyl Violet B, I 3—8—11 Alcohol Blue S F C K Alcohol Blue S F C K Alcohol Soluble Eosine tM Alcohol Yellow G K C R Aldehyde Green K C R Aldehyde Green A Alizarine Black M 12 Alizarine Black 4 B C 3 Alizarine Black 6 B	Acid Violet 4 R S	M	3
Acid Yellow R S. D 3 Acid Yellow S (see Naphthol Yellow S) Acid Yellow S (see Naphthol Yellow S) Acid Yellow S (see Naphthol Yellow S) Acridine Orange Acridine Orange G Acridine Orange R extra L. S.B. Acridine Orange R extra L. S.B.			
Acid Yellow R S. D 3 Acid Yellow S (see Naphthol Yellow S) H 3 Acid Yellow, 8,822 H 3 Acridine Orange L SB 11 Acridine Orange G Bs 11 Acridine Orange R extra L SB 11 Acridine Red B, B B, 3 B L SB Acridine Scarlet R, 2 R, 3 R L L Acridine Yellow L, S B 7-11 Acthyl Blue B F M Acthyle Bosine 0 11 Acthyl Green K B 3-8-11 Acthyl Green A 3-8-11 Acthyl Green A 3-8-11 Akme Yellow L B, I 3-8-11 Acthyl Violet B, I 3-8 All Alcohol Blue B, Br S, By, C R, D, K L, L, R D Alcohol Blue S F C K K Alcohol Soluble Eosine tM Alcohol Yellow G K C R Aldohol Yellow G K C Aldazine Pink B I 2 Aldehyde Green Alizarine Black 4 B C 3 Alizarine Black 4 B C 3 Alizarine Black 6 B C 3			3
Acid Yellow S (see Naphthol Yellow S)			
Acid Yellow, 8,522			3
Acridine Orange L. S. B. 11—7 Acridine Orange G. Bs. 11 Acridine Orange N. O. L. Ac. Acridine Orange R. extra. L. S. B. 1 Acridine Red B. B. B. 3. B. L. S. B. 11 Acridine Scarlet R. 2. R. 3. R. L. L. Acridine Yellow L. S. B. 7—11 Acthyl Blue B. F. M. M. Acthylene Blue O. 11 Acthylene Blue O. 11 Acthyl Green K. B. 3—8—11 Acthyl Green A. 3—8—11 Acthyl Violet B. I. 3—8 Alcohol Blue B. B. S. By, C. R. D., K. L. L. R. D. Alcohol Blue B. B. S. By, C. R. D., K. L. L. R. D. Alcohol Soluble Eosine t. Alcohol Yellow G. K. C. R. Alcohol Yellow R. K. Aldehyde Green M. 12 Alizarine Black M. 12 Alizarine Black B. C. 3 Alizarine			
Acridine Orange G Bs 11 Acridine Orange N O L Acridine Orange R extra L, S B 11 Acridine Red B, B B, 3 B L, S B 11 Acridine Scarlet R, 2 R, 3 R L L Acridine Yellow L, S B 7—11 Aethyl Blue B F M M Aethylene Blue O 11 Aethyl Eosine S 3—8—11 Aethyl Green A 3—8—11 Aethyl Violet B, I 3—8 Alcohol Blue B, Br S, By, C R, D, K L, L, R D Alcohol Blue B, Br S, By, C R, D, K L, L, R D Alcohol Soluble Eosine tM Alcohol Yellow G K, C R Alcohol Yellow G K, C R Alcohol Yellow R K Aldehyde Green K Alizarine Black M 12 Alizarine Black B C 3 Alizarine Black B C 3			. 3
Acridine Orange N O. L Acridine Orange R extra L S B Acridine Red B, B B, 3 B. L S B Acridine Scarlet R, 2 R, 3 R L Acridine Yellow L S B 7-11 Aethyl Blue B F M Aethyle Bosine Aethyl Green A 3-8-11 Aethyl Violet B, I 3-8-11 Akme Yellow L 3-8 Alcohol Blue B, Br S. By, C R, D, K L, L, R D D Alcohol Blue S F C K K Alcohol Soluble Eosine tM Alcohol Yellow G K, C R Alcohol Yellow G K, C R K Alcohol Yellow R K K Aldehyde Green Alizarine Black M 12 Alizarine Black 4 B C 3 Alizarine Black 6 B C 3	Acridine Orange	L, S B	11 - 7
Acridine Orange N O. L Acridine Orange R extra L S B Acridine Red B, B B, 3 B. L S B Acridine Scarlet R, 2 R, 3 R L Acridine Yellow L S B 7-11 Aethyl Blue B F M Aethyle Bosine Aethyl Green A 3-8-11 Aethyl Violet B, I 3-8-11 Akme Yellow L 3-8 Alcohol Blue B, Br S. By, C R, D, K L, L, R D D Alcohol Blue S F C K K Alcohol Soluble Eosine tM Alcohol Yellow G K, C R Alcohol Yellow G K, C R K Alcohol Yellow R K K Aldehyde Green Alizarine Black M 12 Alizarine Black 4 B C 3 Alizarine Black 6 B C 3	Acridine Orange G	Bs	11
Acridine Red B, B B, 3 B. L, S B Acridine Scarlet R, 2 R, 3 R. L Acridine Yellow L, S B Aethyl Blue B F. M Aethylene Blue O Aethyl Eosine 3-8-11 Aethyl Green A 3-8-11 Aethyl Violet B. I 3-8-11 Akme Yellow L 3-8 Alcohol Blue B, Br S. By, C R, D, K L, L, R D Alcohol Blue S F C Alcohol Soluble Eosine tM Alcohol Yellow G K, C R Alcohol Yellow G K K Aldazine Pink B I 2 Aldehyde Green M 12 Alizarine Black M 12 Alizarine Black C 3 Alizarine Black C 3	Acridine Orange N O	L	
Acridine Scarlet R, 2 R, 3 R L Acridine Yellow L, S B 7—11 Aethyl Blue B F M M Aethylene Blue O 11 Aethyl Fosine 3—8—11 Aethyl Green A 3—8—11 Aethyl Violet B. I 3—8—11 Akme Yellow L 3—8 Alcohol Blue B, Br S. By, C R, D, K L, L, R D Alcohol Blue S F C K Alcohol Eosine tM Alcohol Soluble Eosine tM Alcohol Yellow G K, C R Aldazine Pink B I 2 Aldehyde Green K Alizarine Black M 12 Alizarine Black 4 B C 3 Alizarine Black 6 B C 3	Acridine Orange R extra	L, S B	11
Acridine Yellow L, S B 7—11 Aethyl Blue B F M Aethylene Blue O 11 Aethyl Eosine			
Aethyl Blue B F. M Aethylene Blue O 11 Aethyl Eosine			
Aethylene Blue O 11 Aethyl Eosine K B 3-8-11 Aethyl Green A 3-8-11 Aethyl Violet B. I 3-8-11 Akme Yellow L 3-8 Alcohol Blue B, Br S, By, C R, D, K L, L, R D N Alcohol Blue S F C K Alcohol Bolue K C R K Alcohol Soluble Eosine tM K Alcohol Yellow G K C R K Aldazine Pink B I 2 Aldehyde Green K 12 Alizarine Black M 12 Alizarine Black C 3 Alizarine Black C 3	Acridine Yellow	L, S B	7-11
Aethylene Blue O 11 Aethyl Eosine K B 3-8-11 Aethyl Green A 3-8-11 Aethyl Violet B. I 3-8-11 Akme Yellow L 3-8 Alcohol Blue B, Br S, By, C R, D, K L, L, R D N Alcohol Blue S F C K Alcohol Bolue K C R K Alcohol Soluble Eosine tM K Alcohol Yellow G K C R K Aldazine Pink B I 2 Aldehyde Green K 12 Alizarine Black M 12 Alizarine Black C 3 Alizarine Black C 3	Aethyl Blue B F	M	
Aethyl Eosine K B 3—8—11 Aethyl Green A 3—8—11 Aethyl Violet B I 3—8—11 Akme Yellow L 3—8 Alcohol Blue B, Br S, By, C R, D, K L, L, R D Alcohol Blue S F C K Alcohol Soluble Eosine tM Alcohol Soluble Eosine tM Alcohol Yellow G K, C R Aldazine Pink B I 2 Aldazine Black M 12 Alizarine Black 4 B C 3 Alizarine Black 6 B C 3	Aethylene Blue	0	11
Aethyl Green K B 3—8—11 Aethyl Green A 3—8—11 Aethyl Violet B. I 3—8—11 Akme Yellow L 3—8 Alcohol Blue B, Br S. By, C R, D, K L, L, R D Alcohol Blue S F C K Alcohol Eosine tM Alcohol Soluble Eosine tM Alcohol Yellow G K, C R Aldazine Pink B I 2 Aldazine Pink B I 2 Alizarine Black M 12 Alizarine Black 4 B C 3 Alizarine Black 6 B C 3	Aethyl Eosine		
Aethyl Green .A 3—8—11 Aethyl Violet .B. I 3—8—11 Akme Yellow .L .B. I 3—8 Alcohol Blue .B. Br S. By, C R, D, K L, L, R D Alcohol Blue S F C .K. K Alcohol Soluble Eosine .tM Alcohol Yellow G .K. C R <td></td> <td></td> <td>811</td>			811
Aethyl Violet B. I 3—8—11 Akme Yellow L 3—8 Alcohol Blue B, Br S, By, C R, D, K L, L, R D K Alcohol Blue S F C K K Alcohol Eosine tM M Alcohol Soluble Eosine tM K Alcohol Yellow G K, C R K Alcohol Yellow R K K Aldazine Pink B I 2 Aldehyde Green M 12 Alizarine Black M 12 Alizarine Black 4 B C 3 Alizarine Black 6 B C 3			-8-11
Akme Yellow L 3—8 Alcohol Blue B, Br S, By, C R, D, K L, L, R D K Alcohol Blue S F C K K Alcohol Eosine tM K Alcohol Soluble Eosine tM K Alcohol Yellow G K, C R K Alcohol Yellow R K K Aldazine Pink B I 2 Aldehyde Green M 12 Alizarine Black M 12 Alizarine Black 4 B C 3 Alizarine Black 6 B C 3			-S11
Alcohol Blue			3-8
Alcohol Blue S F C. K Alcohol Eosine tM Alcohol Soluble Eosine tM Alcohol Yellow G K, C R Alcohol Yellow R K Aldazine Pink B I 2 Aldehyde Green M 12 Alizarine Black 4 B C 3 Alizarine Black 6 B C 3	Alcohol Blue B. Br S. Bv, C R. D. K L	, L, R D	
Alcohol Eosine tM Alcohol Soluble Eosine tM Alcohol Yellow G K. C. R. Alcohol Yellow R K. Aldazine Pink B I. 2 Aldehyde Green Alizarine Black M. 12 Alizarine Black 4 B C. 3 Alizarine Black 6 B C. 3	Alcohol Blue S F C	K	
Alcohol Soluble Eosine tM Alcohol Yellow G K, C R Alcohol Yellow R K Aldazine Pink B I 2 Aldehyde Green M 12 Alizarine Black M 12 Alizarine Black 4 B C 3 Alizarine Black 6 B C 3	Alcohol Eosine	tM	
Alcohol Yellow G K, C R Alcohol Yellow R K Aldazine Pink B I 2 Aldehyde Green M 12 Alizarine Black M 12 Alizarine Black 4 B C 3 Alizarine Black 6 B C 3	Alcohol Soluble Eosine	tM	
Alcohol Yellow R K Aldazine Pink B I 2 Aldehyde Green M 12 Alizarine Black M 12 Alizarine Black 4 B C 3 Alizarine Black 6 B C 3	Alcohol Yellow G	K.CR	
Aldazine Pink B I 2 Aldehyde Green Alizarine Black Alizarine Black 4 B Alizarine Black 6 B			
Aldehyde Green Alizarine Black Alizarine Black 4 B Alizarine Black 6 B C 3			2
Alizarine Black M 12 Alizarine Black 4 B C 3 Alizarine Black 6 B C 3			
Alizarine Black 4 B			12
Alizarine Black 6 B	Alizarine Black 4 B	C	
Alizarine Black D RM			

Name of Dye. Agent.	Method.
Alizarine Black R, patented	
Alizarine Black R, patentedC M	3
Alizarine Black R T	12
Alizarine Black S	
Alizarine Black S, patented	4
Alizarine Black S, in paste	
Alizarine Black S R A	4
Alizarine Black S W, in paste	3 4
Alizarine Black T, patented	
Alizarine Blue	
Alizarine Blue in paste	
Alizarine Blue A	
Alizarine Blue A BB A Co	
Alizarine Blue B A P M	
Alizarine Blue B R 3 GBy	
Alizarine Blue C GBy	
Alizarine Blue C R	
Alizarine Blue C S	
Alizarine Blue D B	
Alizarine Blue D G	
Alizarine Blue D N W	
Alizarine Blue D R 8	
Alizarine Blue G B A C	
Alizarine Blue G S	
Alizarine Blue G WBy	
Alizarine Blue J GSch	
Alizarine Blue NP K	
Alizarine Blue N A extraP I	ζ 4
Alizarine Blue N BP B	4
Alizarine Blue N G patented P K	4
Alizarine Blue N G G powderP K	4
Alizarine Blue O D R	t
Alizarine Blue RBy, M	1 49
Alizarine Blue R R	
Alizarine Blue S powder	[4
Alizarine Blue S pasteB, By, M	1 4
Alizarine Blue soluble powder A B SB A Co) 4
Alizarine Blue S A PBy	
Alizarine Blue S C A	
Alizarine 1,400 T	
Alizarine Blue Black S W	3
Alizarine Bordeaux B in paste 20%By	7 - 4
Alizarine Bordeaux B D in pasteBy	4
Alizarine Bordeaux C	[4
Alizarine Bordeaux GBy	
Alizarine Bordeaux G GBy	
Alizarine BrownB, B	
Alizarine Brown in powder	I 13
Alizarine Brown in paste	[13
Alizarine Brown A S	
Alizarine Brown G NBy	y 4

Name of Dye,	Agent.	Method.
Alizarine Brown O D R		4
Alizarine Brown R	A P M	4
Alizarine Brown R B	Bv	4
Alizarine Brown S O	P K	4
Alizarine Cardinal	Bv	6b
Alizarine Carmine	B A Co	4
Alizarine Carmine Blue B and G	By	4
Alizarine C A	B A Co	4
Alizarine Cyanine G in paste	By	4
Alizarine Cyanine Black G in paste	By	4
Alizarine Cyanine R in paste		4
Alizarine Cyanine R A extra		4
Alizarine Cyanine 2 R		4
Alizarine Cyanine 3 R		4
Alizarine Dark Blue		4
Alizarine Dark Blue S		4
Alizarine Fast Red		3
Alizarine for Violet	Gau	4-6b
Alizarine G D		
Alizarine G I	B	17b
Alizarine Green B	D	4
Alizarine Green C	By	4
Alizarine Green C E paste	By	4
Alizarine Green C G	By	4
Alizarine Green C K	By	4
Alizarine Green E B	Bs	4
Alizarine Green G	D	4
Alizarine Green, paste	Р К	4
Alizarine Green S in paste	B	4
Alizarine Grenat R		6Ն
Alizarine Indigo S in paste	B	4
Alizarine Lanacyl Blue B B	C	2a
Alizarine Lanacyl Blue 3 B	C	2a
Alizarine Lanacyl Navy Blue B patented.	C	2a
Alizarine Lanacyl Blue R		2a
Alizarine Lanacyl Violet B, patented		2a
Alizarine Maroon paste		6b
Alizarine No. 1	N	4
Alizarine No. 1 B	M	4-6b
Alizarine No. 6	M	6b
Alizarine No. 10	N	6b
Alizarine Olive O D	At	4
Alizarine Orange	Gau	
Alizarine Orange A in paste	В	6b
Alizarine Orange A O	B A Co	6b
Alizarine Orange A O P	B A Co	6Ъ
Alizarine Orange G	By, M	6b
Alizarine Orange N	M	6b
Alizarine Orange powder	M	4-6b
Alizarine Orange R	By	6b
Alizarine P	B A Co	

Name of Dye. Agent.	Method.
Alizarine Powder S A B A Co	4—6b
Alizarine Powder WBy	4—6b
Alizarine Red	4
Alizarine Red S	6b
Alizarine R G	17
Alizarine R X	4
Alizarine SaphirolBy	3
Alizarine Saphirol BBy	3-4
Alizarine S C	
Alizarine Scarlet, patented	17 3
Alizarine S D G	
Alizarine S X	17b
Alizarine S X extraBy	17b
Alizarine VI	17b
Alizarine Violet	
	4-9-17
Alizarine Violet extraAt	4
Alizarine Violet paste	4
Alizarine Violet, 1007 TC	4
Alizarine Viridine pasteBy	4
Alizarine W S powder	4—6b
Alizarine XBy	17a
Alizarine Yellow	4
Alizarine Yellow A pasteB	17
Alizarine Yellow A WB	4
Alizarine Yellow C pasteB	17
Alizarine Yellow F.S	4
Alizarine Yellow G G	4
Alizarine Yellow G G W powder	4
Alizarine Yellow L WP K	4
Alizarine Yellow O DAt	4
Alizarine Yellow paste	4
Alizarine Yellow RBy, C R, M	4
Alizarine Yellow R W powder	4
Alkali BlueA, B, Br S, By, C, C R, D, D H, F, G, J,	
K, L, Lev, M, N I, O, P, P S, Sch, tM,	
Alkali Blue 6B	•
Alkali Blue D	
Alkali Blue X GBr S	
Alkali Brown	12
Alkali Brown RL. P	· 12
Alkali GreenBr S	
Alkali Red	2 2
Alkali Violet	23
Alkali Violet RBy	
Alkali YellowĎ	
Alkali Yellow R	
Alpha-Naphthol Orange	
Alpine Bleu	3
Alsace GrayFi	
Alsace GreenF T M	
Alsace Green JF T M	

Name of Dye. Agent.	Method.
Amaranth B K, C, F, Lev, M, M Lv, P, R D	
Amaranth extraP	3
Amethyst Violet K	
Anil Blue R	
Anileïne	
Aniline Brown	3-11
Aniline Green	0
Aniline Orange	
Aniline Pink	11
Aniline Purple	
Aniline Red K B	3-8-11
Aniline Violet	0 0 11
Aniline Yellow D H	
Aniline Yellow extra	12
Anisidine Ponceau	14
Anisoline	3-8-11
Anisol Red	0 0 11
Anthracene Acid Black L W	3
Anthracene Acid Black S F	3
Anthracene Acid Black S T	3
Anthracene Acid Brown B	3-4-5
Anthracene Acid Brown G	3-4-5
Anthracene Acid Brown N	3-4-5
Anthracene Acid Brown R	3-4-5
Anthracene Acid Brown S W patentedC	3
Anthracene Black	•
Anthracene Blue S	4
Anthracene Blue S W X	•
Anthracene Blue W BB	6b
Anthracene Blue W GB	
Anthracene Blue W RB	4
Anthracene Brown paste	13
Anthracene Brown G pasteBv	. 13
Anthracene Brown R pasteBy	13
Anthracene Croceïne B. G	
Anthracene Dark BlueB	
Anthracene Green (see Coeruleine)	4-9-13
Anthracene RedJ. By	3-4
Anthracene Red BSch	3
Anthracene Scarlet O RSch	3
Anthracene Violet	4-9-13
Anthracene Yellow pasteBv	4
Anthracene Yellow B N	4
Anthracene Yellow C	3-4-5
Anthracene Yellow G G	4
Anthracene Yellow RC	4
Anthracite Black BC	3
Anthragallol (see Anthracene Brown)	13
Anthramine YellowAt	4
Apollo RedG	3
Archil Red 3 V NS S	3

	Agent.	Method.
Arnica Yellow	G	12
Arsen Fuchsine		3-8-11
Atlas Orange	Br S	3-8
Atlas Red	Br S	
Atlas Searlet		3
Auramine GB.		2-7-11
Auramine O, I, II, concB		2-7-11
Arabian Black		12b
Aurantiaxxxxxxx		8
Aureoline		12—12b
Aureosine		
AurineGr, Lo, L P, Mo.	, R D	
Aurine R		et.
Aurotine		Gb
Azaleïne		
Azaline		
Azamine 4 B (see Benzo Purpurine 4 B).	N	8
Azarine R	M	8
Azarine S	M	
Azin Blue alcohol soluble		
Azin Green G B	IX	11
Azin Green G. B	1.	3
Azin Green S	M	7-11
Azo Acid Black B		3
Azo Acid Black G	M.	3
Azo Acid Black R	M	3
Azo Acid Blue B	M	3
Azo Acid Blue 4 B	By	3
Azo Acid Brown	Re	3
Azo Acid Fuchsine B	М	3
Azo Acid Fuchsine G	M	3
Azo Acid Ruby	D	3
Azo Acid Ruby 2 BD.	вк	3
Azo Acid Violet 4 R	By	3
Azo Acid Yellow	Å	3-8
Azo Black O	M	3
Azo Black-Blue	O	12b
Azo Blue	, Lev	12e
Azo Bordeaux	By	3
Azo Bordeaux	()	
Azo Brown O	M	3
Azo Brown V	ss	3
Azo Carmine B	B	3
Azo Carmine G paste	B	3
Azo Cardinal G	A	
Azo Chromine	G	4
Azo Coccine 7 B (see Cloth Red G. By)	$\dots \Lambda$	
Azo Coccine G (see Tropacoline 0000).		
Azo Coccine 2 R	A	3
Azo Cockineal	By	3

Name of Dye.	Agent.	Method.
Azo Coralline	D	3
Azo Corinth		12e
Azo Crimson L	By	3
Azo Crimson S	By	3
Azo Diphenyl Blue		· ·
Azo Eosine	Bv	3
Azo FlavineB,	D tM	3-8
Azo Fuchsine Acide	RF	4
Azo Fuchsine B	Bv	3
Azo Fuchsine G	Bv	3
Azo Fuchsine G N		3
Azo Fuchsine S		3
Azo Galleïne		4
Azo Green	Bv	4
Azo Grenadine S	Bv	3
Azo Mauve B	Ò	12e
Azo Mauve R		12e
Azo Orange R	D H	12
Azo Orseille R		
Azo Orseilline		
Azo Orcelleïne		
Azophenine Blue G	N J	
Azophenine Blue R		
Azophor Black S Printing	M	
Azophor Blue D	M	15b
Azophor Orange Printing	M	15b
Azophor Red P N	M	15b
Azorceine		
Azo Red A		
Azo Rubyt	M, Lev	3
Azo Ruby A	C	3
Azo Ruby S.	A	3
Azo Ruby 2 S	A	
Azo Saffranine	G	3
Azo Turkey Red	0	
Azo Violet	L, Lev	12e
Azo Yellow B K, I	, K, M	3-8
Azo Yellow M		3-8
Azo Benzol Fast Crimson	P K	12
Azuline		
Azurine		
Basle Blue		2—11
Basle Blue B B		
Basle Blue R paste		
Basle Blue S		3—8
Bavarian Blue alcohol soluble	A	
Bavarian Blue D B F	A	7—11
Bavarian Blue D S F		
Bengal Blue		
Bengal PinkC	I, DH	3

Name of Dye, Agent,	Method.
Bengaline Blue BK	11
Benzal Green	11
Benzidine Blue	
Benzidine Red	
Benzindamine	
Benzo Azurine GA, By, L, Lev	12e
Benzo Azurine 3 G	12e
Benzo Azurine RBy	12
Benzo BlackBy	12
Benzo Black Blue GBy	12
Benzo Black Blue 5 GBy	12
Benzo Black Blue RBy	12e
Benzo Black BrownBy	12
Benzo Blue B BBy	12
Benzo Blue 3 BBy	12
Benzo Blue B XBy	12b
Benzo Brown BBy	12
Benzo Brown GBy	12
Benzo Brown 5 RBy	12
Benzo Chrome Black BlueBy	12
Benzo Chrome Black BBy	12
Benzo Chrome Black Blue BBy	12
Benzo Chrome Brown BBy	12
Benzo Chrome Brown 5 GBy	12b
Benzo Chrome Brown 3 RBy	12
Benzo Cyanine BBy	
Benzo Cyanine 3 BBy	
Benzo Cyanine RBy	
Benzo Dark BrownBy	
Benzo Fast GrayBy	
Benzo Flavine	9-11
Benzo Flavine No. 2	9-11
Benzo GrayBy	12
Benzo Green GBy	12
Benzo Indigo BlueBy	12
Benzo Nitrol Brown G and 2 RBy	12b
Benzo Nitrol Brown NBy	12
Benzo OliveBy	12
Benzo Orange R	4-12
Benzo Pure Blue	12b
Benzo Purpurine B	12e
Benzo Purpurine 4 B	12e
Benzo Purpurine 6 B A, By, L, Lev	12e
Benzo Purpurine 10 B	12e
Benzo Sky BlueBy	12
Benzoyl Green	0 0
Benzyl Violet	3-8-11
Best Magenta Crystals K B	
Best Violet (Brilliant India Dye)	
Biebrieh Acid Blue	3
Biebrich Acid Blue G patentedK	3

Name of Dye.	Agent.	Method.
Biebrich Acid Blue G G	K	3
Biebrich Acid Blue G G Biebrich Alizarine Black 4 B N patented	К	3
Biebrich Patent Black A N	K	3
Biebrich Patent Black 4 A N		3
Biebrich Patent Black A O		3
Biebrich Patent Black 4 B N		3
Biebrich Patent Black R O	K	3
Biebrich Patent Jet Black		3a
Biebrich Patent Jet Black 3 B O	IX	3
Riabrich Scarlet	1	3
Biebrich Scarlet	CDF	J.
	M. W	3-11
Bismarck Brown		3-11
Bismarck Brown-Yellow Shade		
Bismarck Brown B		11
		11
Bismarck Brown G		3—11
Bismarek Brown T		3-11
Bitter Almond Oil Green	K B	3-8-11
Blackley Blue	Lev	8-11
Bleu alcalin	M LV	
Bleu alcalin 4 B		
Bleu alcalin B V S Ia No. 57	JD	
Bleu belge	O	11
Bleu brillant Diamine G		20 1
Bleu Chromazone		3-4
Bleu Coupier à l'alcool		11
Bleu de Bengale	D ₁	11
Bleu de ciel alcalin	115	6b
Bleu de ciel soluble		s-11
Bleu de Lyons	V	3-8-11
Bleu de nuit		3-3-11
Bleu de Paris		
Bleu Diamine B B		
Bleu Diamine B B B	M In	
Bleu Diamine B X	MIL	
Bleu Diamine 3 R		
Bleu Diamine R W	MIL	
Bleu direct		
Bleu en pâte	N J	
Bleu fluorescent	TS	
Bleu glacier	T	3-8-11
Bleu lumière	р	0 11
Bleu Marin		3-11
Bleu métaphenylène B		0-11
Bleu méthyl	MLv	s-11
Bleu méthylène nouveau N	M L.	0 11
Bleu méthylène nouveau N G G	M Ly	
Bleu neutre	MLv	
Bleu noir Diamine B		
Bleu noir Diphenyle	Ğ	12
Died non Diphentic		

Name of Dyc. Agent.	Method.
Bleu solide R, 3 R	
Bleu soluble purD W, L, M Ly	811
Bleu nouveau Diamine G	
Bleu nouveau Diamine R	
Bleu nouveau Diamine P	
Bleu opale	
Bleu opale	
Bleu pour impression	
Bleu pur à l'alcool	
Bleu pur Diamine	
Bleu pur Diamine F F M Ly	
Blue asozin	3
Blue alcohol soluble	3
Blue B J B	2
Dive Dieds D	3
Blue Black B	3
Blue Black No. 1	
Blue Black No. 2Bai	
Blue Black 5534H	3
Blue B SC, P	. 3
Blue 3 B SP	. 3
Blue C B alcohol soluble D H	
Blue C B water soluble D H	
Blue extraR D	7 - 11
Blue Green SB	
Blue 77 G A C	12f
Blue Rougeant P P W Printing	
Blue water soluble for wool and silk J	8-11
Bordeaux BA, B K, Lev, L P, M, R F	3-8
Bordeaux B LC	3-8
Bordeaux B XBy	3
Bordeaux C O VÅ	3-12f
Bordeaux D H D H	3-8
Bordeaux Diamine B	3-8
Bordeaux Diamine S M Ly	3-8
Bordeaux extraBy	3-12f
Bordeaux G	3—8
Bordeaux GBy	3
Bordeaux R extra	3—8
Bordeaux S	3—8
Bottle Green Beh	
Brazeline	10
	12
	4
BrexalineAt	
Brexaline At Brahma Orange Z	4
Brexaline At Brahma Orange Z Brahma Red B, B B. Z	4
Brexaline At Brahma Orange Z Brahma Red B, B B. Z Brahma Red 6 B Z	4
Brexaline At Brahma Orange Z Brahma Red B, B B Z Brahma Red 6 B Z Bright Yellow T B, L	12
Brexaline At Brahma Orange Z Brahma Red B, B B. Z Brahma Red 6 B Z Bright Yellow T B, L Brilliant Alizarine Blue D paste By	12 4
Brexaline At Brahma Orange Z Brahma Red B, B B Z Brahma Red 6 B Z Bright Yellow T B, L Brilliant Alizarine Blue D paste By Brilliant Alizarine Blue, G, R By	12
Brexaline At Brahma Orange Z Brahma Red B, B B. Z Brahma Red 6 B. Z Bright Yellow T. B, L Brilliant Alizarine Blue D paste By Brilliant Alizarine Blue, G, R. By Brilliant Alizarine Blue D paste By	12 4
Brexaline At Brahma Orange Z Brahma Red B, B B Z Brahma Red 6 B Z Bright Yellow T B, L Brilliant Alizarine Blue D paste By Brilliant Alizarine Blue, G, R By	12 4

Name of Dye. Agent.	Method.
Brilliant Azurine 5 G	12
Brilliant Black BB	3
Brilliant Blue	
Brilliant Bordeaux S	3
Brilliant Carmoisine 0	
Brilliant Cochineal 2 R	
Brilliant Cochineal 4 R	
Brilliant Chrome Red pasteBy	
Brilliant Congo G	12e
Brilliant Congo R	12e
Brilliant Cotton Blue greenishBy	11
Brilliant CroceineBy	3812
Brilliant Croceine bluish	
Brilliant Croceine 3 BBy	
Brilliant Croceine 9 B	_ 3
Brilliant Croceine M	
Brilliant Cyanine Blue RBy	4
Brilliant Diazine Blue B	
Brilliant Diazine Blue B B	
Brilliant Geranine B patentedBy	12
Brilliant Geranine 3 BBy	
Brilliant Green. B, By, C, C J, C R, D H, F, K, K B,	
M, M Ly, N I, O, P S, R E, tM	3—8—11
Brilliant Green crystalsKchl	3
Brilliant India Dye	
Brilliant Opaline	
Brilliant Orange GM	3
Brilliant Orange G	
Brilliant Orange O	3
Brilliant Orange R	3
Brilliant Orcelleine, patented	3
Brilliant Orseille C	
Brilliant Ponceau GB K, C, Lev	3
Brilliant Ponceau G G	3—8
Brilliant Ponceau 4 RBy	
Brilliant Ponceau 5 RBy, C, D	3
Brilliant Purpurine	
Brilliant Purpurine R	3
Brilliant Purpurine 5 B	
Brilliant RedSch	3
Brilliant Red DI	3
Brilliant Red Congo G	3
Brilliant Red 2038 T	12
Brilliant Saffranine G	11
Brilliant ScarletLev	. 3
Brilliant Scarlet 6 R	3
Brilliant Sulfon Azurine RBy	2
Brilliant YellowSch	3 3—8
Brilliant YellowtM	3—8 12e
Brilliant Yellow	3-8
Brilliant Yellow SB	35

Name of Dye.	Agent.	Method.
Bronze Diamine G	M Lv	12—15
Brun au chrome		10 10
Brun Bismarek No. 259		
Brun Bismarek E E	M Ly	
Brun Congo G	R-F	12
Brun Corinthe B		12e
Brun Corinthe G		12e
Brun Diamine R	\dots M Ly	12 & 15e
Brun Diamine 3 G	M Ly	12 & 15e
Brun Diamine M	M Ly	12 & 15e
Brun Diamine V	M Ly	12 & 15
Brun Diphényle		108
Brun Mikado 3 G O	L, S B	12f 12f
Brun Mikado M	T. S. D.	121 12f
Brun Naphtine a		1.01
Brun Naphtine 3		4
Brun P M		•
Brown B B X		12
Brown J E		2
Brown J E E E	P	3
Brown N		3
Brown S D M		12
Brown S D P		12
Buffalo Rubin		.3
Butter Yellow	• • • • • • • • •	
Caehou de Laval	Р	15
Cachou de Laval S		15
Cachou Diamine	M Ly	12
Campanuline		11
Canarine		
Canelle		3-11
Capri Blue G N		11
Capri Green B		11
Capri Green G	<u>.</u> <u>.</u>	
Capri Green G G	b	12b
Carbazol Yellow W		120
Cardinal		1.
Cardinal Red S		
Carminenaphte		
Carminenaphte J		
Carminenaphte Grenat	В Н	
Carmoisine	B, By, SS	3
Carmoisine conc		3
Carnotine		12—12b
Caronbier	В Н	
Catechu Brown F K		
Catechu Brown F D K		
Catechy Brown G K		

Name of Dye. Agent.	Method.
Catechu 2 G K	
Cattu Italiano L D	15
Celeste Blue B	
Cerasine	
Cerasine Orange G	
Cerasine Red	
CeriseB. C. D H, K B, P, P S, N J	
Cerotine Orange C extra	
Cettonine	3
Chicago Blue BA	
Chicago Blue 4 B	
Chicago Blue 6 BA	
Chicago Blue R	
Chicago Blue 2 R	
Chicago Blue 4 R	
Chicago Blue R W	
Chicago GrayG	
Chicago Orange	
China Blue	8—11
China Green Crystals K B.	
Chinoline Blue	
Chinoline GreenB	
Chinoline Red	0 6
Chinoline Yellow	38
Chinoline Yellow alcohol soluble B. By, R. F.	12
Chloramine Brown C	12
Chloramine OrangeBy Chloramine YellowBy	
Chlorhydrate of Toluidine	17
Chlorophenine G	11
Chlorophenine Orange A A	2
Chlorophenine Orange R	2
Chlorophenine R	3-7-12
Chlorophenine Y	3-7-12
Chromazone Red	3
Chromanil Brown G G	12
Chromanil Brown RA	12
Chromanil Black R F	
Chrome BlueBy	4
Chrome Blue 2 B	4
Chrome Blue 4 B	4
Chrome Blue B N	4
Chrome Blue No. 470	4
Chrome Blue P E	4
Chrome Blue R	4
Chrome Blue 2 R	4
Chrome Blue 3 R	4
Chrome BordeauxBy	4
Chrome Brown G	4
Chrome Brown R O	3
Onfome rast black b	3

Name of Dyc. Apent.	Method.
Chrome Fast Yellow GA	
Chrome Fast Yellow 2 G	
Chrome Fast Yellow R	
Chrome GreenBy	4
Chrome OrangeBy	4
Chrome PruneBy	4
Chrome Red pasteBy	4
Chrome Ruby PasteBy	4
Chrome VioletBy	
Chrome Violet powderBy, G	4
Chronie Violet pasteBy	4
Chrome Yellow	
Chrome Yellow DBy	4
Chrome Yellow P	4
Chromine GK	12
Chroniogene I	3b
Chromotrope 2 B	3b
Chromotrope 6 B	36
Chromotroje 8 BM	3b
Chromotrope 10 B	3b
Chromotrope F B	3b
Chromotrope 2 R	3
Chromotrope S	3b
Chromotrope S N	3h
Chromotrope S R	3b
ChrysamineBy	12
Chrysamine G	12e
Chrysamine G GBy	12
Chrysamine R	12e
Chrysaniline	
Chrysaureïne	
Chryseoline	
Chrysoidine eryst M	2a-7-11
Chrysoidine eryst	
+11 11'	1-7-11
Chrysoidine G	1-7-11
Chrysoidine J	1-7-11
Chrysoidine R	1-7-11
Chrysoidine R	1-1-11
Chrysoidine R	11
Chrysoidine Y Lev	1-7-11
Chrysoidine C E E	1-1-11
Chrysoine	3
Chrysoïne	.)
Chrysoline	
Chrysophenine	19.5
Chrysophenine G	12e
Cinereine	12
Cinnabar Searlet	
Cinnamine S	
Cinnamon Brown	2 11
Chinamon Brown S	3-11

Name of Dye. Agent. Method.
Citronine Fi, S B 3—8 Citronine A L C Citronine A C 3—8 Citronine A A C 3—8 Citronine A H E P 3 Citronine N E P 3 Clayton Aurotine Cl O 4 Clayton Carnotine Cl O 3—8 Clayton Cloth Red Cl O 3 Clayton Yellow Cl O 12e Clematine G 11 Cloth Brown reddish By, L 4 Cloth Brown yellowish By, L 4 Cloth Red M 4 Cloth Red B By 4 Cloth Red B A A 4 Cloth Red G By 4 Cloth Red G By 4 Cloth Red G extra By 4 Cloth Red G extra By 4 Cloth Red G oxtra By 4 Cloth Red G oxtra By 4 Cloth Red G oxtra
Citronine Br S Citronine A L Citronine A C Citronine 2 A extra S S 7 Citronine A H E P Citronine N E P Clayton Aurotine Cl O 4 Clayton Carnotine Cl O 3 Clayton Cloth Red Cl O 3 Clayton Red Cl O 3 Clayton Yellow Cl O 12e Clematine G 11 Cloth Brown reddish By, L 4 Cloth Brown yellowish By, L 4 Cloth Red M 4 Cloth Red B D 4 Cloth Red B D 4 Cloth Red B A A 4 Cloth Red G By Cloth Red G By Cloth Red G cextra By
Citronine A L Citronine A C 3—8 Citronine 2 A extra S S 7 Citronine A H E P 3 Citronine N E P 3 Citronine N E P 3 Clayton Aurotine Cl O 4 Clayton Cloth Red Cl O 3 Clayton Cloth Red Cl O 3-8 Clayton Red Cl O 3-8 Clayton Yellow Cl O 3 Clayton Yellow Cl O 12e Clematine G 11 Cloth Brown reddish By, L 4 Cloth Brown yellowish By, L 4 Cloth Red B M 4 Cloth Red B By, D 4 Cloth Red B By, D 4 Cloth Red B A 4 Cloth Red G By 4-9 Cloth Red G O 4 Cloth Red G O 4 Cloth Red G O <td< td=""></td<>
Citronine A C 3—8 Citronine 2 A extra S S 7 Citronine A H E P 3 Citronine N E P 3 Clayton Aurotine Cl O 4 Clayton Carnotine Cl O 3 Clayton Cloth Red Cl O 3-8 Clayton Red Cl O 3 Clayton Yellow Cl O 12e Clematine G 11 Cloth Brown reddish By, L 4 Cloth Brown yellowish By, L 4 Cloth Red M 4 Cloth Red By, L 4 Cloth Red B By, D 4 Cloth Red B By, D 4 Cloth Red B A 4 Cloth Red G By 4 Cloth Red G O 4 Cloth Red G extra By 4 Cloth Red G extra By 4
Citronine 2 A extra S S 7 Citronine A H E P 3 Citronine N E P 3 Clayton Aurotine Cl 0 4 Clayton Carnotine Cl 0 3 Clayton Cloth Red Cl 0 3-8 Clayton Red Cl 0 3 Clayton Yellow Cl 0 12e Clematine G 11 Cloth Brown reddish By, L 4 Cloth Brown yellowish By, L 4 Cloth Red M 4 Cloth Red B By, D 4 Cloth Red B O 4 Cloth Red B A A 4 Cloth Red G By 4-9 Cloth Red G By 4 Cloth Red G extra By 4 Cloth Red G extra By 4 Cloth Red 3 G extra By 4
Citronine A H E P 3 Citronine N E P 3 Clayton Aurotine Cl O 4 Clayton Carnotine Cl O 3 Clayton Cloth Red Cl O 3-8 Clayton Red Cl O 3 Clayton Yellow Cl O 12e Clematine G 11 Cloth Brown reddish By, L 4 Cloth Brown yellowish By, L 4 Cloth Red M 4 Cloth Red B By 4 Cloth Red B By 4 Cloth Red B A A 4 Cloth Red G By 4 Cloth Red G By 4 Cloth Red G extra By 4 Cloth Red G extra By 4 Cloth Red G extra By 4
Citronine N E P 3 Clayton Aurotine Cl 0 4 Clayton Carnotine Cl 0 3 Clayton Cloth Red Cl 0 3-8 Clayton Red Cl 0 3 Clayton Yellow Cl 0 12e Clematine G 11 Cloth Brown reddish By, L 4 Cloth Brown yellowish By, L 4 Cloth Orange By, L 4 Cloth Red M 4 Cloth Red B By, D 4 Cloth Red B A 4 Cloth Red B A 4 Cloth Red G By 4-9 Cloth Red G By 4 Cloth Red G O 4 Cloth Red G extra By 4 Cloth Red G extra By 4
Clayton Aurotine Cl 0 4 Clayton Carnotine Cl 0 3 Clayton Cloth Red Cl 0 3-8 Clayton Red Cl 0 3 Clayton Yellow Cl 0 12e Clematine G 11 Cloth Brown reddish By, L 4 Cloth Brown yellowish By, L 4 Cloth Orange By, L 4 Cloth Red M 4 Cloth Red B By, D 4 Cloth Red B O 4 Cloth Red B A 4 Cloth Red B By 4-9 Cloth Red G By 4 Cloth Red G O 4 Cloth Red G extra By 4 Cloth Red G extra By 4
Clayton Carnotine Cl 0 3 Clayton Cloth Red Cl 0 3-8 Clayton Red Cl 0 3 Clayton Yellow Cl 0 12e Clematine G 11 Cloth Brown reddish By, L 4 Cloth Brown yellowish By, L 4 Cloth Orange By, L 4 Cloth Red M 4 Cloth Red B By, D 4 Cloth Red B O 4 Cloth Red B A A 4 Cloth Red G By Cloth Red G Cloth Red G O 4 Cloth Red G extra By 4 Cloth Red G extra By 4 Cloth Red G extra By 4
Clayton Cloth Red Cl O 3—8 Clayton Red Cl O 3 Clayton Yellow Cl O 12e Clematine G 11 Cloth Brown reddish By, L 4 Cloth Brown yellowish By, L 4 Cloth Red M 4 Cloth Red B By, D 4 Cloth Red B O 4 Cloth Red G By 4—9 Cloth Red G By 4 Cloth Red G O 4 Cloth Red G extra By 4 Cloth Red G extra By 4 Cloth Red G extra By 4
Clayton Red Cl 0 3 Clayton Yellow Cl 0 12e Clematine G 11 Cloth Brown reddish By, L 4 Cloth Brown yellowish By, L 4 Cloth Orange By, L 4 Cloth Red M 4 Cloth Red B By, D 4 Cloth Red B O 4 Cloth Red B A A 4 Cloth Red G By 4-9 Cloth Red G By 6 Cloth Red G O 4 Cloth Red G extra By 4 Cloth Red G extra By 4 Cloth Red G extra By 4
Clayton Yellow Cl O 12e Clematine G 11 Cloth Brown reddish By, L 4 Cloth Brown yellowish By, L 4 Cloth Orange By, L 4 Cloth Red M 4 Cloth Red B By, D 4 Cloth Red B O 4 Cloth Red B A A 4 Cloth Red G By 4-9 Cloth Red G By 4 Cloth Red G extra By 4 Cloth Red G extra By 4 Cloth Red 3 G extra By 4
Cloth Brown reddish By, L 4 Cloth Brown yellowish By, L 4 Cloth Orange By, L 4 Cloth Red M 4 Cloth Red B By, D 4 Cloth Red B O 4 Cloth Red B A 4 Cloth Red B By 4-9 Cloth Red G By 4 Cloth Red G O 4 Cloth Red G extra By 4 Cloth Red 3 G extra By 4
Cloth Brown yellowish By, L 4 Cloth Orange By, L 4 Cloth Red M 4 Cloth Red B By, D 4 Cloth Red B O 4 Cloth Red B A A 4 Cloth Red G By 4-9 Cloth Red G By Cloth Red G Cloth Red G extra By 4
Cloth Brown yellowish By, L 4 Cloth Orange By, L 4 Cloth Red M 4 Cloth Red B By, D 4 Cloth Red B O 4 Cloth Red B A A 4 Cloth Red G By 4-9 Cloth Red G By Cloth Red G Cloth Red G extra By 4
Cloth Red M 4 Cloth Red B By, D 4 Cloth Red B 0 4 Cloth Red B A 4 Cloth Red B By 4-9 Cloth Red G By 4 Cloth Red G O 4 Cloth Red G extra By 4 Cloth Red 3 G extra By 4
Cloth Red B By, D 4 Cloth Red B 0 4 Cloth Red B A A 4 Cloth Red 3 B extra By 4-9 Cloth Red G By 0 4 Cloth Red G extra By 4 Cloth Red G extra By 4 Cloth Red 3 G extra By 4
Cloth Red B .0 4 Cloth Red B A .A 4 Cloth Red 3 B extra By 4-9 Cloth Red G By Cloth Red G extra By 4 Cloth Red G extra By 4 Cloth Red 3 G extra By 4
Cloth Red B A A 4 Cloth Red 3 B extra By 4-9 Cloth Red G By Cloth Red G 4 Cloth Red G extra By 4 Cloth Red 3 G extra By 4
Cloth Red 3 B extra By 4—9 Cloth Red G By Cloth Red G 0 4 Cloth Red G extra By 4 Cloth Red 3 G extra By 4
Cloth Red G By Cloth Red G O 4 Cloth Red G extra By 4 Cloth Red 3 G extra By 4
Cloth Red G O 4 Cloth Red G extra By 4 Cloth Red 3 G extra By 4
Cloth Red G extra By 4 Cloth Red 3 G extra By 4
Cloth Red 3 G extraBy 4
Cloth Pod C A
Cloth Red 3 G A
Cloth Red O 4
Cloth Red O B 3
Cloth Red O B, F R B O
Cloth Red R
Cloth Scarlet G K 3 Cloth Scarlet R K 3
Coccein 3 B
CoccininM
Coccinin B
Cochenille brillant 2 R
Cochenille brillant 4 R
Cochineal Red A
Cochineal Scarlet GSch 3
Cochineal Scarlet P SBy
Cochineal Scarlet 2 RSch 3
Cochineal Scarlet 4 RSch 3
Cochineal Substitute
Coeruleïne paste
Coeruleine A paste
Coeruleine B paste
Coeruleine B W paste
Coeruleine S powder or pasteB, By, D H, Fi, M 4-9-13.

Name of Dye.	Agent,	Method.
Coeruleïne S W powder or paste		4-9-13
Columbia Black B		4-5-15
Columbia Black B B	Α	
Columbia Black F B	Α	2a.
Columbia Black R	AA.	411
Columbia Blue G		
Columbia Blue R		
Columbia Brown R	ΑΑ.	12
Columbia Chrome Black B B	Λ	2a
Columbia Fast Blue 2 G	A	Zil
Columbia Green		
Columbia Red 8 B	AA.	
Columbia Yellow		
Congo	T Tou	1 10
Congo B		1—12
Congo Blue 3 B		10
Congo Blue B X		12 12b
Congo Blue 2 B X	1 D.	12 12
Congo Brown G	A Tow	12
Congo Brown R	A, Lev	12
Congo Corinth B	Or T Tor	
Congo Corinth G	Dr. I. Lev	12e 12e
		12e
Congo Fast Blue B		
Congo Fast Blue R Congo G R	A D.	100
Congo Orange R	A, DV	12e 12
Congo Orange G		12
Congo P		
Congo Pure Blue		12b
Congo 4 R		· 12e
Congo Red		12—12f
Congo Ruby		12-121
Congo Violet		3—12f
Congo Yellow en pâte	A Dy	3—121
Coralline		
Coralline (red)		
Coralline (yellow)		
Coreïne		4
Coreïne A B		4
Coreïne A R		4
Coreïne R R		4
Cotton Black B		12b
Cotton Blue		1.0
Cotton Blue		8-11
Cotton Blue 3 B		12g
Cotton Blue 6 B extra		8—11
Cotton Blue G		12g
Cotton Blue R		11
Cotton Bordeaux		
Cotton Brown 3 G, R		

Name of Dye.	Agent.	Method.
Cotton Brown	В	
Cotton Brown N	C	12
Cotton Brown R		4
Cotton Orange G		12
Cotton Orange R	В	12
Cotton Ponceau		
Cotton Red		12
Cotton Red 4 B		12e
Cotton Scarlet		3-8-176
Cotton Scarlet 3 B conc		
Cotton Yellow G		12
Cotton Yellow G B	d	12 12b
Coupiers Blue		120
Cresotine Yellow G		12
Cresotine Yellow R		12
Cresyl Blue B B		1-
Cresyl Blue 6B		11
Cresyl Blue B B S O	Bs	11
Cresyl Blue R R	Bs	11
Cresyl Fast Violet B, B B		
Croceïne A Z	C	
Croceïne B		3
Croceine 3 B		38
Croceine 3 B X		3
Crocéine brillante M	M Ly	
Croceine OrangeBy, B K, K,	Lev. R F.	3
Croceine Scarlet 3 B	By, K	3—12g
Croceine Scarlet 4 B X		3
Croceïne Scarlet 7 B		3
Croceine Searlet 8 B		3
Croceine Scarlet 10 B		3
Croceine Scarlet O extra		3—8
Cross Dye Black		12a
Cross Dye Drab		12a 12
Crumpsall Direct Fast Brown B		12
Crumpsall Direct Fast Brown M	Lar	12
Crumpsall Direct Fast Brown O	Ler	12
Crumpsall Fast Yellow Y Y F D	Lev	3
Crumpsall Yellow	Lev	3-4
Crystal Ponceau		3
Crystal Ponceau 6 R	M. M Lv	3
Crystal Violet	B	3-8-11
Crystal Violet 5 B O	I	3-8-11
Crystal Violet O	B	3-8-11
Cumassi Black		3c
Cumassi Mauve Blue		3c
Cumidine Ponceau		3
Cumidine Red		3
Curcumeine	A, B K	3

Name of Dye. Agent.	Method,
Curcumine	3-8
Curcumine S A Br I C D	3-8-14
Curcumine Substitute	3
CurcuphenineCl O	9
Cutch Brown D	2a-7-11
Cutch Brown G G	12b
Cyanine	14.,
Cyanine B M	3
Cyanol extra	3
Cvanosine	Ü
Cyanosine alcohol soluble K. M.	
Cyanosine alcohol soluble K, M Cyklamine Mo	2-7
	~ .
Dahlia	3-8-11
Dark Blue B L	3
Dark Brown M	2a—7—11
Dark Green B	6d
Dark Green 682 Br S	12
Deep Wool Black 2 B	2
Deep Wool Black 3 B	2
Delphin Blue	~
Delta Purpurine 5 B	12e
Delta Purpurine 7 B	190
Delta Purpurine GBy	12e
Diamant Gros Cristany M Ly	126
Diamine Azo Black B patented	12a-
Diamine Azo Blue No. 5 154 natented C.	1~0
Diamine Azo Blue No. 72,122 patented	3—12e
Diamine Azo Blue R patented	12e
Diamine Azo Blue R R patented	12e
Diamine Black B	15
Diamine Black B H	15
Diamine Black B O	15
Diamine Black H W	15
Diamine Black R O	15
Diamine Black-Blue B	10
Diamine Blue B	12 or 15
Diamine Blue B B	12-15
Diamine Blue 3 B	12-15
Diamine Blue B G patented	10
Diamine Blue B X C M Ly	12
Diamine Blue N C	12b
Diamine Blue 3 R C M Ly	12-15
Diamine Blue R W	12-15
Diamine Blue S R X patented	12
Diamine Blue No. 50 patented	1.5
Diamine Blue No. 52 patented	
Diamine Blue No. 53 patented	
Diamine Blue No. 55 patented	
Diamine Blue 72.918	12
Diamine Blue Black E	12-15
	12-13

Name of Dye.	Agent.	Method.
Diamine Blue Black R	C	12-15
Diamine Bordeaux B	C	
Diamine Bordeaux S	C	
Diamine Brilliant Blue G		12
Diamine Bronze C	C	12—15e
Diamine Bronze G	M Ly	12-15
Diamine Brown A	C	12—15e
Diamine Brown B	C	12—15e
Diamine Brown 3 G	C	12—15e
Diamine Brown M		12—15e
Diamine Brown V	M Ly	12-15
Diamine Brown Q Q	C	12—15e
Diamine Brown 3,833	C	12—15e
Diamine Brown No. 30a patented	C	
Diamine Brown No. 31 patented	C	
Diamine Brown No. 32 patented	C	
Diamine Brown No. 33 patented	C	*-
Diamine Brown No. 34 patented	C	
Diamine Brown No. 35 patented		
Diamine Brown No. 36 patented	C	-
Diamine Brown No. 37 patented	C	
Diamine Brown 2 O patented	C	
Diamine Catechine B patented	C	12-15
Diamine Catechine G patented	C	12-15
Diamine Catechu	C	12
Diamine Dark Blue B	C	12
Diamine Deep Black Cr patented	C	12a
Diamine Deep Black 0000	C	12-15
Diamine Deep Black R B	C	12—15
Diamine Deep Black S S		12-15
Diamine Deep Dark Blue B and R	C	12c
Diamine Fast Brown G		12
Diamine Fast Red		3-12
Diamine Fast Red F	M Lv	3
Diamine Fast Yellow A		12f
Diamine Fast Yellow B		12b
Diamine Fast Yellow R		12
Diamine Flavine R	$\mathbf{B}_{\mathbf{y}}$	12
Diamine Gold	C	12
Diamine Gold Yellow	M Ly	12a
Diamine Gray G	C	12
Diamine Green B		12
Diamine Green G		12
Diamine New Blue G		12
Diamine New Blue R		12
Diamine Nitrazol Black B		12
Diamine Nitrazol Brown B patented		12-15
Diamine Nitrazol Brown B D patented		12-15
Diamine Nitrazol Brown G patented		12-15
Diamine Nitrazol Brown R D patented		12—15
Diaminogene B	M Ly	12

Name of Dye.	Agent.	Method.
Diaminogene extraC,	M Ly	
Diaminogene B R Diaminogene Blue B B	C	
Diaminogene Blue B B	C	12
Diaminogene Blue G	С	12
Diamineral Black B		12
Diamineral Blue R patented		12b
Diamine Orange B	C	12
Diamine Orange D	C	12
Diamine Orange G	C	12
Diamine Orange G C	C	12
Diamine Pink B D, B extra	C	12
Diamine Pure BlueC,		12b
Diamine Pure Blue F F		
Diamine Red B	C	12e
Diamine Red 3 B		12e
Diamine Red 10 B		12
Diamine Red N OC,		12a
Diamine Scarlet B		3—8
Diamine Scarlet 3 B		3—8
Diamine Sky Blue	C	12
Diamine Sky Blue F F	С	12
Diamine Steel Blue L		12a
Diamine Violet N		12
Diamine W O		12
Diamine Yellow N		12
Diamine Yéllow paste	C	12
Diamine Yellow R paste		12
Diamond Black		4
Diamond Brilliant Blue G		
Diamond Brown		
Diamond Fast Acid Black		12
Diamond Fast Blue		12
Diamond Fast Red		12
Diamond Fast Yellow		12
Diamond Flavine G		4
Diamond Green		4
Diamond Green		11
Diamond Green B		3-8-11
Diamond Green G	В	3-8-11
Diamond Green erystals	KB	
Diamond Orange paste		
Diamond Yellow G paste		4
Diamond Yellow R paste		4
Dianil Black		4
Dianil Black G paste	M	4
Dianil Black R paste		4
Dianil Blue B		4
Dianil Blue G		4
Dianil Blue R		4
Dianil Blue Black E		4
Dianisidine Blue	By, M	

Name of Dye.	Agent,	Method.
Diphenyl Blue E F S		12
Diphenyl Blue N T	G	12
Diphenyl Blue Black	G	12
Diphenyl Brown	G	12
Diphenyl Brown B	G	12
Diphenyl Brown R		12
Diphenyl Brown R R	G	12
Diphenyl citronine G		12e
Diphenyl Fast Black	G	• 12
Diphenyl Fast Yellow G	G	12
Diphenyl Indigo Blue		12
Diphenyl Orange		.12
Diphenyl Red		12
Diphenyl Violet R		12
Diphenyl Yellow		12
Direct Black B	S S	12
Direct Black B F G	A	12
Direct Black G B N		12
Direct Black K		12b
Direct Black R		12
Direct Black X		12
Direct Blue		
Direct Blue 5093		12
Direct Blue B		12
Direct Blue B		
Direct Blue 2 B		12
Direct Blue 3 B X		
Direct Blue 5 B		12
Direct Blue G	Sch	12
Direct Blue G 97		12
Direct Blue O	B L	12
Direct Blue R	I	12
Direct Blue 2 R	S S	12
Direct Blue Black B	By	
Direct Brown 5,092	S W	12
Direct Brown B B	Bs	12
Direct Brown G G	By	12
Direct Brown G X	\dots Bs	12
Direct Brown J		12
Direct Brown M		12
Direct Brown N X		12b
Direct Brown R		12
Direct Brown S		12
Direct Brown S D P		12
Direct Brown V X		12b
Direct Brown 130		3
Direct Brown 131		3
Direct Catechu Brown		
Direct Deep Black E extra		12
Direct Deep Black R	\dots By	12b
Direct Deep Black R W	\dots By	12

Name of Dye.	* Apent.	Method.
Direct Deep Black T		100000
Direct Fast Brown B		12
Direct Fast Brown G G		3
Direct Gray		2
Direct Gray B	7	12
Direct Gray B		12
Direct Grav Reddish	N.I	1-
Direct Green		12b
Direct Green B		12
Direct Green B C		12
Direct Green B X		12
Direct Green Y		3
Direct Green Y Y C	S W	12
Direct Indigo Blue A	I	12b
Direct Indigo Blue B K		12b
Direct Lemon Yellow		12
Direct Orange	Fi	
Direct Orange G 69	Sch	12
Direct Orange R	S S	12
Direct Orange 2 R	K	8—12f
Direct Orange Y	S S	12
Direct Oriol Yellow	G	12
Direct Red		12
Direct Red		
Direct Red		
Direct Red B		12b
Direct Red E T		12
Direct Red No. 55		12b
Direct Scarlet B conc		12f
Direct Scarlet G		12f
Direct Scarlet R		12f
Direct Union Black		12
Direct Yellow		
Direct Yellow		
Direct Yellow		12f
Direct Yellow		3
Direct Yellow C		12
Direct Yellow G		12f
Direct Yellow 2 G		12f
Direct Yellow 3 G		12f
Direct Yellow R		12
Direct Yellow T		3
Double Brilliant Scarlet G		3
Double Brilliant Scarlet 3 R		3
Double Green S F		•
Double Ponceau 2 R		3
Double Ponceau 3 R. Double Ponceau 4 R.		3
Double Scarlet		3
Double Scarlet extra S		3
Double Scarlet G		3
Dodoic Scatter G		3

N	A	Method.
Name of Dye. Double Searlet R	Agent.	Method.
		3
Double Searlet 2 R	tM	3
Eboli Blue	Pe.	2—12
Eboli Blue B		12f
Eboli Green B		12e
Ecarlate B	р	3
Eearlate brillante		3
Ecarlate Crocéïne 3 B		3
Eearlate d'Eosine B		3-8
Ecarlate Diamine B		2-3-7
Ecarlate Diamine 3 B		2-3-7
Ecarlate J		3—7
Ecarlate J J	Mo	37
Eearlate V		37
Echurine		
Eclipse Red		12e
Emerald Green erystals		3-7-11
Emin Red		3
Empire Orange G	Beh	12
English Brown		3—11
Eosamine B		
Eosine	C J, F	3—8
Eosine A	B	38
Eosine B		3—8
Eosine B B	I	
Eosine 10 B	C, M Ly	
Eosine B N	B	3—8
Eosine B W		3-8
Eosine bleu		
Eosine bleuâtre		3—17b
Eosine bluish		3—8—12 f
Eosine D H		3 - 8 - 12f
Eosine D H V		3—8—12 f
Eosine extra		
Eosine G extra		
Eosine G G F		3—8
Eosine J		3—17
Eosine 3 J		3—8
Eosine 4 J extra		3-8
Eosine J J F		3—8 3—8
Eosine K S ord		3—8
Eosine S		2 0
Eosine Scarlet B		3—8 3—8—12f
		3-8-121
Eosine alcohol soluble		
Eosine water soluble		3 -8
Eosine water soluble		3-8
Erie Blue G G	DI S, A	38
Erika B	A DE	3-12
IIIIRG D		3-12

Name of Dye. Agent.	Method.
Erika GLev	
Erika 4 G N	12b
Eriocyanine	3
Erioglaucine	3
Erythrine XB	S
Erythrobenzine	
ErythrosineB. Br S, CJ, DH, F, M, tM	3—11
Erythrosine A	3
Erythrosine B	
Erythrosine D	
Erythrosine G	
Erythrine X	3
Essaine	- 33
Ethyl Purple 6 B P K	3
Eupitton Acid	3
Excelsior Lake Scarlet J N	3
Excelsion Black	3
Extra Superfine (fine) Brilliant Green CrystalsK B	9
Extra cuprimit (mic) Diminut often civitais I D	
Fast Acid Blue BBv	
Fast Acid Blue R	3—8
Fast Acid Fuchsine B	•
Fast Acid Green B	
Fast Acid Green B S	3
Fast Acid Green B Z	12
Fast Acid Ponceau	
Fast Acid Red A	
Fast Acid Red B	3
Fast Acid Scarlet D H, S	
Fast Acid Violet A 2 R	3
Fast Acid Violet B	3-8
Fast Acid Violet 10 BBy	3
Fast Acid Violet R	
Fast Azo Granat	
Fast BlackL	11
Fast Black BB	12b
Fast Black B SB	
Fast BlueBr S	3
Fast Blue B	3
Fast Blue 2 B for Cotton	11
Fast Blue 6 B for Wool	
Fast Blue alcohol solubleA, B K	
Fast Blue for Cotton	
Fast Blue greenish	3
Fast Blue Black pasteL	11
Fast Blue Black M pasteL	11
Fast Blue GBdt	4 2
Fast Blue R	3
Fast Blue 3 R	11
rast blue is for cotton, in crystals	11

. Name of Dye. Agent.	Method.
Fast Blue III R	11
Fast Bordeaux O	4
Fast BrownBy	3
Fast Brown 3 B	3
Fast Brown G	3
Fast Brown N	3
Fast Brown O N T yellowish	3-8
Fast Brown RB L	12
Fast Brown 25	3
Fast Claret	3
Fast Cotton Blue B	11
Fast Cotton Blue R	- 11
Fast Cotton Blue R R	11
Fast Cotton Blue 3 R crystals	11
Fast Cotton Brown R	12
Fast Cotton Orange 6 R extraBs	12
Fast Cotton Yellow 6 G extraBs	12
Fast Cotton Yellow OBs	12
Fast Cotton Yellow RBs	12
Fast Direct Blue BBdt	12
Fast Direct Blue GBs	12
Fast Direct Brown B BBs	12
Fast Direct Brown GBs	12
Fast Fulling Blue R R	4
Fast Green	6a
Fast Green	3-8-11
Fast GreenBy	3
Fast Green extraBy	3
Fast Green extra bluishBy	3
Fast Green B	
Fast Indigo Blue R	3
Fast Light GreenBy	3
Fast Mordant Yellow G	11
Fast Navy Blue	11
Fast Navy Blue G M, R MK	11
Fast Navy Blue M M, R MK	11
Fast Neutral Violet B	11
Fast New Blue for Cotton	11
Fast Pink for Silk D H	-
Fast Ponceau BB	3
Fast Ponceau 2 B	3
Fast Red	3
Fast RedF, L	3
Fast Red AA, B, B K, By, C R, K, Lev, tM	3
Fast Red BB, B K	3-8
Fast Red BN J	
Fast Red B TBy, D H, Lev	3
Fast Red CB	3
Fast Red DB	3-8
Fast Red E	3

Name of Dye. Agent.	Method.
Fast Red E BB	38
Fast Red 7 B	0 0
Fast Red N SBy	3—8
Fast Red OM	3
Fast Red RA C	3
Fast Red R YB K	3
Fast Red S	3
Fast Scarlet	12e
Fast ScarletK	3
Fast Scarlet BK	38
Fast Silk Gray OM	
Fast Silk Red cf. Fast Red 7 B N J	
Fast Violet	
Fast Violet bluishBy	• 3
Fast Violet reddishBy	3
Fast YellowB, By, L P, M Ly, S B	
Fast YellowBr S	
Fast YellowB	3-8
Fast YellowtM	3
Fast Yellow GB K, D H, K	
Fast Yellow greenish	
Fast Yellow RB K, K	3—8
Fast Yellow S	
Fast Yellow 4 SS S	3
Fat Ponceau	
Fine Blue (see Aniline Blue alcohol soluble).	
Fine New Green Crystals K B	
Fine Violet	3—8—11
Flavaniline	3—5—11
Flavaniline S	
Flavaurine	
Flavazol	3
Flaveosine	J
Flavinduline	11
Flavophenine	
Fluoresceïne	38
Fluorescent I, II, III, IVII	
Fluorescent Blue	
Plugninding	
ridorindine	
Fluorindine	
Fond RougeCz	3
Fond Rouge Cz Formyl Violet 4 B C	3 3
Fond Rouge Cz Formyl Violet 4 B. C Formyl Violet 6 B. C Formyl Violet 8 B patented. C	
Fond Rouge Cz Formyl Violet 4 B. C Formyl Violet 6 B. C Formyl Violet 8 B patented. C Formyl Violet 10 B. C	3
Fond Rouge Cz Formyl Violet 4 B. C Formyl Violet 6 B. C Formyl Violet 8 B patented. C Formyl Violet 10 B. C Formyl Violet S 4 B. C, M Ly	3
Fond Rouge Cz Formyl Violet 4 B. C Formyl Violet 6 B. C Formyl Violet 8 B patented. C Formyl Violet 10 B. C Formyl Violet S 4 B. C, M Ly Formyl Violet S 5 B. C	3 3 3
Fond Rouge Cz Formyl Violet 4 B.	3 3 3 3
Fond Rouge Cz Formyl Violet 4 B. C Formyl Violet 6 B. C Formyl Violet 8 B patented. C Formyl Violet 10 B. C Formyl Violet S 4 B. C, M Ly Formyl Violet S 5 B. C Fuchsia I Fuchsiacine I	3 3 3 3 3a 11
Fond Rouge Cz Formyl Violet 4 B.	3 3 3 3a

Name of Dye. Agent,	Method.
Fuchsine acide	
Fuchsine B O O	2
Fuehsine J O O	2
Fuchsine S B D S P S	2-8
Fuehsine Searlet	2 0
Fulling BlueK	4
runing blue G	3
Fulling Blue R	3
Fulling Green	0
Fulling Green S	
Fulling Red B	'
Fulling Red B	
Fulling Red F G G	
Fulling Red F R	
Fulling Red G	
Fulling Red R	
Fulling Yellow	
Fulling Yellow O O	4
Tuning Tenow O OC	
Gallamine Blue	
Gallamine Blue pasto	4
Gallanil Green	
Gallanil Green	4
Gallanil Indigo P	4
Gallarin A Gallarin A Gallarin A	4
Gallazin A D H	4
Gallein paste	4 - 9 - 13
Galleïn A paste	4
Gallein W powder	4
Gallocyanine B S D H	4-13
Gallocyanine D II	4-13
Gallocyanine paste D	3
Galloeyanine pasteBy	4
Galloflavine in paste	413
Gambine	6a
Gambine B	46a
Gambine G paste	6a
Gambine R paste	6a
Gambine Y paste	6a
Gambine Y D S	6a
Gambine Yellow	6a
uanani i cilow = Martins Yellow D E	3
Gentian Blue 6 R	
Genuanin	11
Genuine VioletK B	
Geranium	11
Geranium G N	3
Geranine B B	
Geranine G	
Germania Red	
inrone Dur	11
Giroffé in paste and powder D H	

Name of Dye.	Agent.	Method.
Gladioline		
Gloria Black B patented	C	3d
Glycine Blue	Ki	
Glycine Corinth	Ki	12e
Glycine Red	Ki	12e
Gold Brown		3-11
Golden Yellow		3
Gold Orange		3-11
Gold OrangeB K,	Br D	3—8
Gold Orange for Cotton	H +M	.,
Gold Yellow		3-8
Granat Brown		0
Gray R & B		
Greenish Blue		
Green Powder = Methyl Green		7
Grenadine	TDS	
Grenat S		
Grenat soluble	36 T	
Gris Diamine G = Diamine Gray G [C]	M LY	12
Gris direct B en poudre	P	11
Gris direct J en poudre	<u>F</u>	11
Gris direct R en pâte	<u>P</u>	11
Gris direct 4 R en pâte	P	11
Gris Diphényle	<u>G</u>	
Gris spécial R en pâte	P	11
Guinea Green B	A	3-8
Guinea Green B V		3-8
Guinea Red 4 R	A	3
Guinea Violet 4 B	A	3
	_	
Half Wool Black		
Half Wool Black S	C	
Harmaline		
Havana Blue W		12f
Helianthine		3—8
Helianthine		38
Heligoland Blue B	N J	
Heligoland Blue G	N J	
Heligoland Blue R	N J	
Heligoland Blue G A	Jy	12
Heligoland Blue 2 R	Jy	12
Heligoland Brown	N J	
Heligoland Yellow	N J	12
Heligoland Red	N J	
Heliochrysine		
HeliotropeA,		2-12
Heliotrope au tannin	M Lv	
Heliotrope BA.	By. L	12e
Heliotrope B	K	11
Heliotrope 2 B	By. L	12
Heliotrope 2 B		11

Name of Dyc. Agent.	Method.
Heliotrope 2 BA	
Helioxanthine = Diphenylamine Orange	3
Helvetia Blue	3-11
Helvetia GreenBi	
Hessian BlueL	
Hessian BordeauxL, S B	12
Hessian Brilliant Purple	12
Hessian Brown B BL	11
Hessian Brown M ML	12
Hessian Purple B	12e
Hessian Purple D	12e
Hessian Purple N	12e
Hessian Violet	12e
Hessian Yellow	12e
Hoechst New Blue	3
Hofmann's Violet K B	3-8-11
Hofmann's Violet N	11
Homophosphine GL	11
HydroleïneR E	11
Hydroleïne Induline	
Hydroleïne Marine RR E	
Hydroleïne Primula B	
Transition of the state of the	
Immedial Black V extra patented	12
Imperial BlackAt	3
Imperial Green Crystals K B	
Imperial Red Violet	3
Imperial SearletBy	3
Imperial Violet CrystalsAt	3
Indamine Blue N B	12
Indamine Gray N J	
Indamine 3 R	11
Indamine 6 R	11
Indamine T D	**
Indazine M	11
Indazurine	
Indian YellowBy	3-8
Indian Yellow G	0 11
Indian Yellow R	12
Indigene D, FBy	1~
Indigo Blue N	3
Indigo Blue R BS S	3
Indigo Blue S G N	3
Indigo Powder 1,006	3
Indigo SaltK	3
Indisine	
Indoehronine T	4
Iodoïne Blue	*
Indol Blue R	11
Indophenine extraBy	12
Indophenol powderD H	15
independent portret in the second sec	

- Name of Dye. Agent.	Method.
Indophenol White, paste	
IndophorB	
Induline B, B K, By, C J, C R, D, L P, N J, R D, W	3
Induline alcohol soluble. B. B. K. By, C. J. C. R. F. I.	
LP, NJ, RD, tM, W	
Induline 3 BL P	3
Induline 6 B	3
Induline B EP	3
Induline N NB	3
Induline R & B	3
Induline Scarlet	11
Marron, Crimson, Brown (see Primuline)	
Intense BlueBv	3
Iodine Eosine B.	3-17
Iodine Eosine G	
Iodine EosineMo	
Iodine Green	
Iodine Violet	3-8-11
Iris BlueB	12
Iris VioletB	3
Isatine Yellow	
Iso Diphenyl Black B	12
Iso Diphenyl Black R	12
Iso Purpurine	4
Iso Rubin	3-8-11
Italian Green	12f
Janus Blue R	
Janus Bordeaux B	3
Janus Brown B	3
Janus Brown R	3
Janus Gray B	3
Janus Grav B B	3
Janus Green B	3
Janus Red B patented	' 3
Janus Yellow RKchl	3
JasmineG	3-8
Jaune II M Ly	
Jaune acide D H, L P	
Jaune acide	
Jaune acide C	
Jaune Anglais	
Jaune Anthracène C=Anthracene Yellow C [C] M Ly	3-4
Jaune Anthracène $GG = Anthracene Yellow GG[C]$	
M Ly	4
Jaune brillant L P	
Jaune brillant	
i Apilipo Vellor	
Jaune d'Aniline = Aniline Yellow Orange IV.	
(Orange 1 .,	

Name of Dye. Agent.	Method
Jaune Diamine N MLy	
Jaune de Quinoleïne solubleA, B, By, R F	
Jaune direct	
Jaune d'or	
Jaune d'Orient	
Janne G (Matanil extra)	
Jaune Indien G M Ly	
Jaune G (Métanil extra) M Ly Jaune Indien G M Ly Jaune métanile A, B, B K, By, D, D H, Fi, G, K, O,	
S B. tM	
Jaune métanile broméP	
Jaune NaphtholS B	
Jaune resistant au savonP	
Jaune Soleil	
Jaune solideB, By, L P, M Ly, S B	
Jaune solide Diamine A=Diamine Fast Yellow A[C]	
M Ly	
Jaune solide Diamine B = Diamine Fast Yellow B[C]	
M Ly	
Jaune solide N	0 2
Jet Black RBy	2-3
Kaiser Red	3—11
Kaiser Yellow	3—11
Kanarine D H	J
Kanthosine J	
Kanthosine R	
Katigene Black Brown NBy	12e
Kermesine Orange	3
Keton Blue 4 B N solution	3-8
Keton Blue G	3-8
Keton Blue R	3—8
Kresotine Yellow G	12
Kresotine Yellow R	12
Kresol RedB	
Lackmoïd	
Lake Scarlet F R	3
Lake Scarlet F R R	3
Lake Scarlet F R R RC	3
Lake Scarlet G GC	3
Lake Scarlet 2 RC	3
Lanacyl Alizarine Violet R	2a
Lanafuchsine S BC	3
Lanafuchsine S GC	3
Lancaster Yellow	
Lauth's Violet	•
Lazuline Blue RBy	3-11
Leather Brown	3-11
Leather Brown	

Name of Dye Agent.	Method.
Leather Brown	
Leather Yellow	
Light BluetM	
Light Blue superfine alcohol soluble	
Light GreenK B	
Light Green S F bluishB	3—8
Light Green S F yellowishB	
London Blue extraBr S	8-11
Lutécienne	
Luteoline	
Lutetienne	
Lyons Red AS S	2-3-7
Lydine	
Madras Blue BF T M	
Madras Blue B	3
Madras Blue G	12
Madras Blue R RAt	3
Madras Blue R	3
Magdala Red	
Magenta	2-7-11
Magenta Base	2-7-11
Magenta large crystals B	3
Magenta No. I	3-11
Mais	3—8—12f
Malachite Green. A, C J, C R, D H, F, K, K B, M, P S.	3-0-121
R E, tM	3-8-11
Malachite Green BB	3-8-11
Malachite Green GB	3-8-11
Malachite Green alcohol soluble	
Malbery Blue	3-8-11
Malta GravP	11
Malta Gray	3-8-11
Manchester Brown E E	3—12
Manchester Brown P S	3-11
Manchester YellowLev, R D	3
Mandarine	3—8
Mandarine G extra	3-8
Mandarine G RA	3
Marron	
Marron SB	3
Mars Red GB	
Martius YellowB K	3
Mauve	
Mauve Dye	
Mauveine	
Mazarine Blue R N S A C	4
Mekon Yellow G D H	12e
Mekon Yellow R D H	12
Meldolas Blue	11
Merino Blue S S	12

Name of Dye. Agent.	Method.
Merino Blue RP	12
Merino BrownS S	12
Metamine Blue BL	11
Metamine Blue GL	11
Metanil Yellow. A, B, B K, By, D, D H, Fi, G, K, O,	
S B tM	3
Metanil extra Yellow G	3
Metaphenylene Blue B	11
Metaphenylene Blue B B	11
Methyl Alkali BlueB, D H, G, K, O, M	
Methyl Blue	8-11
Methyl Blue for Cotton	3-8-11
Methyl Blue for Silk M L B	
Methyl Blue NewG	
Methyl Cotton Blue	8-11
Methyl Diphenylamine Blue	
Methylene BlueP	11
Methylene Blue BB	11
Methylene Blue B B	11
Methylene Blue B B erystals	11
Methylene Blue B B in powder extraA, B, F, M	11
Methylene Blue D, B B extra	11
Methylene Blue in powder I or D, extra D M, A	11
Methylene Blue B G	11
Methylene Blue B G	11
Methylene Green extra yellowish G coneM	12
Methylene Green extra yellow	12
Methylene Violet D H	1~
Methylene Violet R R A	11
Methylene Violet 3 R A	11
Methyl Eosine	3-8
Methyl Eosine	36
Methyl Green	7
Methyl Green	4
Methylindone B, R	. 11
Methyl Violet B, 2 B. A, B, By, B K, C, C J, C R, F,	. 11
K B, M, N J, O, R D, R E, tM	2 0 11
Methyl Violet 6 BA, B K, C, K, M, N J, O, P	3-8-11
Methyl Violet 6 B crystals	3-8-11
Methyl Violet 7 B	1-11
Methyl Violet 3 B O	3-8-11
Methyl Violet B B, 72 O	2
	12
Methyl Violet C 7 P	12
Methyl Violet S 7 BP K	3
Methyl Violet V 3K	3-8-11
Methyl Water Blue	8—11
Metternieht's Green	
Mikado Brown B	12f
Mikado Brown 3 G OL, S B	12 f
Mikado Brown ML, S B	12f
Mikado Gold Yellow 2 GL	12 f

Name of Dye.	Agent	Method.
Mikado Gold Yellow 4 G		12f
Mikado Gold Yellow 6 G		12f
Mikado Gold Yellow 8 G		12f
Mikado Orange G		12
Mikado Orange B		12
Mikado Orange 2 R	L. S B	12
Mikado Orange 3 R	L. S B	12
Mikado Orange 4 R	L S B	12
Mikado Orange 5 R	L, S B.	12
Mikado Yellow		12
Milling Red F R		3
Milling Red G	C	3
Milling Yellow II		3
Milling Yellow O	C	23
Milling Yellow No. 55		3
Mimosa Moline		12
Mordant Yellow		3
Mordant Yellow O	D	4
Murexide		*
Muscarine		11
THE PERSON NAMED IN COLUMN TO SERVICE OF THE PERSON NAMED IN COLUMN TO SERVICE		11
Naccarat	P	3
Nankin in cake		
Naphthal Yellow E S	Bv	
Naphthaline Pink = Magdala Red.		
Naphthaline Red	N J	
Naphthaline Red = Magdala Red.		
Naphthaline Scarlet = Magdala Red.		
Naphthaline Yellow	.C. D. F	3
Naphthamine Blue 2 B patented	<u>K</u>	12
Naphthamine Blue 5 B patented	K	12
Naphthamine Blue B R	K	12
Naphthamine Blue 2 R patented		12
Naphthamine Blue 3 R patented	V	12 12
Naphthamine Indigo Blue 2 R patented	Ε	12
Naphthamine Indigo Blue 2 B patented	K	12
Naphthamine Indigo Blue 5 B patented	К	12
Naphthandoine B B	С	11-17
Naphthazarine Blue B	0	3
Naphthazarine S	0	4-13
Naphthazarine R		
Naphthazine Blue	L, M	3
Naphthazine Blue		
Naphthazine Blue O	Kchl	3
Naphthazurine B		3
Naphthazurine B B		12
Naphthazurine R		
Naphthazurine R E	0	3
Naphthion Red = Orseille Substitute V.		

Name of Dye. Agent.	Method.
Naphthine SP	
Naphthindone B B, patented, M Ly	11
Naphthindone T	11
Naphtho Cyanine E P	
Naphtho RubyBy	3
Nanhthol Black B	3
Naphthol Black 3 B	3
Naphthol Black 4 B	
Naphthol Black 6 BC, D	3
Naphthol Black 12 B	12
Naphthol Black P	3
Naphthol Black 4 R	3
Naphthol Blue B	11
Naphthol Blue DR E	11
Naphthol Blue R	11
Naphthol Blue 2 B	12
Naphthol Blue 2 B P K Naphthol Blue G	12
Naphthol Blue Black	3
Naphthol Green B	6d
Naphthol Orange	3-8
Naphthol Orange	30
a-Naphthol Orange	
β-Naphthol Orange	3
Naphthol Red	3-8
Naphthol Red O	3-5
Naphthol Red SB	
Naphthol Yellow	
Naphthol Yellow SB, B K, By, C, C R, D H, I, Lev.	
M, O, R F O	
M, O, R F O	
M, O, R F O Naphthol Yellow R S	3
M, O, R F O Naphthol Yellow R S Naphthylamine Brown Naphthylamine Pink D H	
M, O, R F O Naphthol Yellow R S Naphthylamine Brown Naphthylamine Pink Naphthylamine Vellow K	3
M, O, R F O Naphthol Yellow R S Naphthylamine Brown Naphthylamine Pink Naphthylamine Vellow K	3
M, O, R F O Naphthol Yellow R S Naphthylamine Brown Naphthylamine Pink Naphthylamine Yellow Naphthyl Blue Naphthyl Blue Naphthyl Blue 2 B B	
M, O, R F O Naphthol Yellow R S Naphthylamine Brown Naphthylamine Pink Naphthylamine Yellow K Naphthyl Blue K Naphthyl Blue 2 B Naphthyl Blue Black N C	3 12
M, O, R F O Naphthol Yellow R S Naphthylamine Brown Naphthylamine Pink Naphthylamine Yellow K Naphthyl Blue K Naphthyl Blue B Naphthyl Blue Black N C Naphthyl Blue Black M N Y C	3
M, O, R F O Naphthol Yellow R S Naphthylamine Brown Naphthylamine Pink Naphthylamine Yellow K Naphthyl Blue K Naphthyl Blue B Naphthyl Blue Black N C Naphthyl Blue Black M N Y C	3 12 3a 11
M, O, R F O Naphthol Yellow R S Naphthylamine Brown Naphthylamine Pink Naphthylamine Yellow Naphthyl Blue Naphthyl Blue 2 B Naphthyl Blue Black N Naphthyl Blue Black M N Y Naphthyl Blue Black M N Y Naphthylene Blue R crystals Naphthylene Bed Sy	3 12 3a
M, O, R F O Naphthol Yellow R S Naphthylamine Brown Naphthylamine Pink Naphthylamine Yellow Naphthyl Blue Naphthyl Blue 2 B Naphthyl Blue Black N Naphthyl Blue Black N Naphthyl Blue Black M N Y Naphthylene Blue R crystals Naphthylene Red Naphthylene Violet	3 12 3a 11
M, O, R F O Naphthol Yellow R S Naphthylamine Brown Naphthylamine Pink Naphthylamine Yellow Naphthyl Blue Naphthyl Blue 2 B Naphthyl Blue Black N Naphthyl Blue Black N Naphthyl Blue Black M N Y Naphthylene Blue R crystals Naphthylene Red Naphthylene Violet	3 12 3a 11
M, O, R F O Naphthol Yellow R S Naphthylamine Brown Naphthylamine Pink Naphthylamine Yellow Naphthyl Blue Naphthyl Blue Naphthyl Blue Black N Naphthyl Blue Black N Naphthyl Blue Black M N Y Naphthylene Blue R crystals Naphthylene Red Naphthylene Violet Naphthyl Violet Naphthyl Violet Narceine D H	3 12 3a 11
M, O, R F O Naphthol Yellow R S Naphthylamine Brown Naphthylamine Pink Naphthylamine Yellow Naphthyl Blue Naphthyl Blue Naphthyl Blue Black N Naphthyl Blue Black N Naphthyl Blue Black M N Y Naphthylene Blue R crystals Naphthylene Red Naphthylene Violet Naphthyl Violet Naphthyl Violet Narceine D H	3 12 3a 11
M, O, R F O Naphthol Yellow R S Naphthylamine Brown Naphthylamine Pink Naphthylamine Yellow Naphthyl Blue Naphthyl Blue 2 B Naphthyl Blue Black N Naphthyl Blue Black N C Naphthyl Blue Black M N Y C Naphthylene Blue R crystals Naphthylene Red Naphthylene Violet Naphthyl Violet Narceine D II Navy Blue Navy Blue B Navy Blue B Navy Blue B Nay Blue B Navy Blue B	3a 12 3a 11 12b
M, O, R F O Naphthol Yellow R S Naphthylamine Brown Naphthylamine Pink Naphthylamine Yellow Naphthyl Blue Naphthyl Blue Naphthyl Blue Black N Naphthyl Blue Black N Naphthyl Blue Black M N Y Naphthylene Blue R crystals Naphthylene Red Naphthylene Violet Naphthyl Violet Naphthyl Violet Naphthyl Violet Navy Blue	3a 12 3a 11 12b
M, O, R F O Naphthol Yellow R S Naphthylamine Brown Naphthylamine Pink Naphthylamine Yellow Naphthyl Blue Naphthyl Blue Naphthyl Blue Black N Naphthyl Blue Black N Naphthyl Blue Black M N Y Naphthylene Blue R crystals Naphthylene Red Naphthylene Violet Naphthyl Violet Naphthyl Violet Naphthyl Violet Navy Blue	3 12 3a 11 12b
M, O, R F O Naphthol Yellow R S Naphthylamine Brown Naphthylamine Pink Naphthylamine Yellow Naphthyl Blue Naphthyl Blue 2 B Naphthyl Blue Black N Naphthyl Blue Black N C Naphthyl Blue Black M N Y C Naphthylene Blue R crystals Naphthylene Red By Naphthylene Violet Naphthyl Violet Naphthyl Violet Narceine D II Navy Blue S W Navy Blue	3 12 3a 11 12b
M, O, R F O Naphthol Yellow R S Naphthylamine Brown Naphthylamine Pink Naphthylamine Yellow Naphthyl Blue Naphthyl Blue Naphthyl Blue Black N Naphthyl Blue Black N Naphthyl Blue Black M N Y Naphthylene Blue R crystals Naphthylene Violet Naphthylene Violet Narceïne Narceïne Nary Blue Navy Blue Navy Blue Navy Blue B	3 12 3a 11 12b
Naphthol Yellow R S Naphthylamine Brown Naphthylamine Pink Naphthylamine Yellow Naphthyl Blue Naphthyl Blue Naphthyl Blue Black N Naphthyl Blue Black N Naphthyl Blue Black M N Y Naphthyl Blue Black M N Y Naphthylene Red Naphthylene Red Naphthylene Violet Naphthyl Violet Narceïne Nayy Blue Navy Blue Navy Blue Navy Blue B Navy Blue C Navy Blue B Navy Blue C Navy Blue B Navy Blue B Navy Blue S Na	3 12 3a 11 12b
M, O, R F O Naphthol Yellow R S Naphthylamine Brown Naphthylamine Pink Naphthylamine Yellow Naphthyl Blue Naphthyl Blue Black Naphthyl Blue Black N Naphthyl Blue Black N Naphthylene Blue R crystals Naphthylene Red Naphthylene Violet Naphthylene Violet Naphthyl Blue Navy Blue Navy Blue Navy Blue B R R R R R R R R R R R R R	3 12 3a 11 12b
Naphthol Yellow R S Naphthylamine Brown Naphthylamine Pink Naphthylamine Yellow Naphthyl Blue Naphthyl Blue Naphthyl Blue Black N Naphthyl Blue Black N Naphthyl Blue Black M N Y Naphthylene Blue R crystals Naphthylene Red Naphthylene Violet Naphthyl Violet Narceine Naphthyl Violet Navy Blue	3 12 3a 11 12b
M, O, R F O Naphthol Yellow R S Naphthylamine Brown Naphthylamine Pink Naphthylamine Yellow Naphthyl Blue Naphthyl Blue Black Naphthyl Blue Black N Naphthyl Blue Black N Naphthylene Blue R crystals Naphthylene Red Naphthylene Violet Naphthylene Violet Naphthyl Blue Navy Blue Navy Blue Navy Blue B R R R R R R R R R R R R R	3 12 3a 11 12b

Name of Dye,	Agent.	Method.
Neutral Gray G	A	
Neutral Red		
Neutral Scarlet	L	
Neutral Violet	C. M Ly	
Neutral Violet O		1-12
New Blue B	C. I	11
New Blue G	tM	11
New Blue RB K. By, C. C.	R. D. I. tM	11
New Coccine	A. M. R F	3
New Coccine R		3
New Cotton Solid Blue		11
New Croceine		3
New Direct Blue B		12
New Fast Black		
New Fast Blue in paste F, H	Bv	
New Fast Gray		•
New Fuchsine		3-8-11
New Gray		11
New Gray		3-12
New Green		3-8-11
New Green		
New Indigo		12
New Metamine Blue M		
New Methylene Blue B B		11
New Methylene Blue G G		11
New Methylene Blue N		11
New Methylene Blue R		11
New Methylene Blue 3 R	C	11
New Methylene Gray B G	M	11
New Patent Blue B	Bv	3
New Patent Blue 4 B		3
New Pink		
New Phosphine G		11
New Red L	K	3
New Solid Green B B		3-8-11
New Solid Green 3 B	I	3-8-11
New Toluvlene Blue B		12
New Toluvlene Brown B		12
New Toluvlene Brown B B		12
New Victoria Black B		
New Victoria Black Blue		
New Victoria Blue R		
New Victoria Green	В. КВ	3-11
New Yellow	ВК	
New Yellow		3
New Yellow L		
New Yellow		3
Nicholson Blue		
Nigramine		11
Night Blue	В, І	
Night Green		-

	gent. Method
Nigrisine	P
Nigrisine J Nigrosine alcohol solubleA, B, B K, C J, D, G, 1	P 11
Nigrosine alcohol solubleA, B, B K, C J, D, G, I	N J, S, R 3
Nigrosine water solubleA, B K, C J, D, D H, F	, G,
K, N J, P S, R,	, S B 3
Nile Blue 2 B	B 11
Nile Blue N N	
Nile Blue Rp-Nitraniline Red (Nitrosamine Red)	B 11
Nitrazine Yellow	0
Nitrobenzole Fuchsin	KB ClO 12
Nitrosamine Red	B 7—15b
Noir à l'alcool	
M	Lv
Noir-Bleu Diamine E	Ly
Noir de Lyons	.Mo
Noir Diamine B	Ly
Noir Diamine M L = Diamond Black H W [C]. M	LLV
Noir Diamine R O	Ly
Noir mi-laine $\Lambda = \text{Half-wool Black } [C], \dots, M$	Ly Ly
Noir mi-laine $S = Half-wool Black S [C]M$	Ly
Noir Naphthol 6 B	Ly Ly
Noir Naphthylamine 6 B = Naphthylamine Bl	lack
Noir Naphthylamine D	Ly
6 B [C] M Noir Naphthylamine D]
Noir Oxy-Diamine S000 = Oxy-Diamine Deep Bl	LiV
S000 [C] M	Ly
Noir pour laine	
Noir Vidal S	P 12
Non Mordant Cotton Blue	3r S .tM
Nyanza Black B	A
Oenanthinine	р н з—8
Old Gold No. 203 powder A 1	P M 12
Old Gold S C	
Oleïne	B L 12
Opal BlueBr	S, C

Name of Dye, Agent,	Method.
Opaline Black 1,602Bch	12
Opaline 2 GGb	
Opaline 5 G	
Opaline RGb	
Oramine Blue RP K	12
OrangetM	3
Orange AL	3-8
Orange IBy, D H, K, M, R D, R F, tM, W	3—8
Orangé No. 1	3—8
Orange H. B. K. C. R. D. H. F. Fl. I. K. Lev. M. M.	•
Ly, PS, RD, RF, tM, W	3—8
Orangé No. 2 P Orange III D H, P, R D, R F, tM, W	38
Orange III D H, P, K D, K F, tM, W	3-8
Orangé No. 3	3
Orange IV. B, B K, D, D H, F, F1, G, K, L P, M, M Ly,	
P, R D, R F, S B	3-8
Orangé No. 4	. 3
Orange au tannin K = Tannin Orange K [C]M Ly	
Orangé Cerasine G	
Orange E N L	2
Orange E N Z	3
Orange extra	3-8
Orange G	3—8
Orange G	3-8
Orange G GB K, C, D, M Ly	3—3
Orangé G G en cristaux	3
Orange G R X	3
Orange G S	3
Orange G TBy	3
Orange MBi	3
Orange M G pasteP	3
Orangé Mikado GL, S B	12
Orangé Mikado R	12
Orangé Mikado 2 RL. S B	12
Orangé Mikado 3 RL, S B	12
Orangé Mikado 4 R L. S B	12
Orangé Mikado 5 RL. S B	12
Orange M N	3
Orange NK	3
Orange NB, I	3
Orange P	3-8
Orange RS	4
Orange RB K, C, D H, I, tM	3
Orange RB	3
Orange R LC	3
Orange R N	3
Orange R R = Orange R $[B]$ Bi	3
Orange R R LC	3
Orange Red $I = Double Brilliant Scarlet G [A]$	3
Orange TK	3

Name of Dye. Agent.	Method.
Orange T A	
Orange Yellow = Orange G	3
Orcelline II R	
Orcelline No. 4	3
Oriol YellowG	12
Orseille Brillante C = Brillant Orseille C [C]M Ly	
Orseille Brown	
Orseille Red AB	3
Orseille Substitute G	
Orseille Substitute N extra	3
Orseille Substitute V	3
Orseille Substitute V C R, Fi, P	3
Orseille Substitute 3 V NP	3
Orseilline B BBy	3
Orchil crimson powderP K	3
Oxamine Blue BB	
Oxamine Blue BF	
Oxamine Blue 4 BF	
Oxamine Blue RF	
Oxamine Blue 3 R	12
Oxamine Blue R XP K	12
Oxamine Blue Black BF	
Oxamine Blue Black R	
Oxamine MaroonB	12
Oxamine RedB	12
Oxamine Scarlet BF	
Oxamine VioletB	12
Oxblood 8,851Bs	12
Oxy-Diamine Black A patentedC	12
Oxy-Diamine Black A M patented	12
Oxy-Diamine Black A T patented	12
Oxy-Diamine Black B G patented	12
Oxy-Diamine Black B M patentedC	12
Oxy-Diamine Black B Z patented	12
Oxy-Diamine Black D patented	12
Oxy-Diamine Black N patented	12
Oxy-Diamine Black N R patented	12
Oxy-Diamine Black N R T patented	12
Oxy-Diamine Black R R patented	12a
Oxy-Diamine Black S O O O patented	12
Oxy-Diamine Black W patented	12
Oxy-Diamine Blue 3 R patented	12
Oxy-Diamine Orange G patented	12
Oxy-Diamine Orange R patented	12
Oxy-Diamine Yellow G G patented	12
Oxyphenine	
Oxyphenine GoldCl Co	
Paeonine	
Palatine BlackB	
Palatine Orange	

Name of Dye.	Agent.	Method.
Palatine Red		3
Palatine Scarlet		3
Paper Scarlet bluish	M	
Para Blue		11
Para Fuchsine		3-8-11
Paramine Brown C & R	C R	12
Paramine Indigo Blue	C R	12—15b
Paramine Navy Blue R	C R	12—15b
Para Nitraniline Red		
Para Phenylene Blue R	D	11
Para Phenylene Violet	D	11
Para Sulferine S	В К	
Paris Green = Methyl Green		
Paris Violet	Р	11
Paris Violet 6 B	R E	11
Paris Violet 7 B	R E	11
Parma R paste	S	4-11
Patent Atlas Red		
Patent Blue A		3
Patent Blue N superfine conc	M	3
Patent Blue V	M	3
Patent Fustic or Fustin O Wood &	Bedford	4
Patent Fustic G	Bedford	4
Patent Green V		3-8
Patent Orange	A	3
Patent Phosphine	I	12
Paten Rock Scarlet	Br S	12b
Pegu Brown G	L	12
Pelikan Blue		
Perkins Violet		
Persian Yellow	G	4
Persulphocyan Yellow		
Phenameine		
Phenamine		
Phenamine Blue B, G, R	В	3
Phenanthrene Red		
Phenethol Red		
Phénicienne		
Phénicin		
Phenix Red A	C	3
Phenocyanine	D H	6a
Phenocyanine B	D H	6a
Phenocyanine D	D н	ha
Phenograpine T B	I	4
Phenocyanine V S	I	4
Phenol Flavine	0	3
Phenol Black S S	Ву	
Phenolphtaleine		
Phenosafranine		
Phenyl Brown		
Phenylene Black	Р	3

Name of Dye.	Agent.	Method.
Phenylene Black paste	P	2
Phenylene Blue	B K	11
Phenylene Brown		3—11
Philadelphia Yellow G	A	
Phlorine	M. A	
Phloxine	он, г, м	
Phloxine Iodine free	В	
Phloxine P	В	
Phloxine T A	Mo	4.4
PhosphineB, Br S, C, M, M Ly, N I, O, 1	, PS, S B	11
Phosphine II		
Phosphine N	X	0- 7 11
Phosphine P	M	2a—1—11
Phosphine nouvelle G = New Phosphine [UJM Ly	. 11
Phthaline Navy Blue	A P M	•
Phtaline Wool Black B	A P M	3
Picryl Orange		•
Picryl Yellow		
Pigment Brown	в	1.1
Pink		11
Pittakal		12
Pluto Black B	ву	12b
Pluto Black G		12b
Pluto Black R	ву	12—15
Polychromine	G	
Polychromine B		12-1.5
Ponceau acide	Dn, s	
Ponceau aux cristaux 6 R	11	3
Ponceau B extra		
Ponceau B O extra		
Ponceau brillant 4 R		
Ponceau d'Orient	L' Lou M	3
Ponceau G	D DE M	3—8
Ponceau 2 G	D, D K, M	3—3
Ponceau 4 G B	A, D N	
Ponceau G R	M Tv	
Ponceau J	MIT	
Ponceau J J		
Ponceau 2 R	M P P E	
Ponceau 2 R S	M, 1, IV P	
Ponceau 3 R B	Λ	
Ponceau 3 R B	D K	
Ponceau 4 R	A B M	
	R R	
Ponceau 5 R		
Ponceau 6 R		
Ponceau 6 R B		
Ponceau 10 R B		
Ponceau R T		
Lonceau R I		

Name of Dye.	Agent.	Method.
Ponceau S extra = Fast Tonceau 2 B [B]	A	3
Ponceau S S extra	A	
Ponceau Y B		
Pourpre		3-8
Prague Alizarine Yellow G	Ki	4-13
Prague Alizarine Yellow R	L':	4—13
Dei		4-13
Primrose		
Primrose à l'alcool D H	рн	
Primrose à l'alcool		
Primrose soluble	D H	3—12b
Primula		3-8-11
PrimulineB, Br S, By, C, G, K, L F	MLv	12-15
Primuline O		12-15
Printing Black for Wool		
Printing Blue	Α	
Printing Blue H		
Delegation Discord		
Printing Blue B		
Printing Blue R		
Propiol Acid		
Prune pure	C	
Pure Blue	B. I. L	8-11
Pure Blue B S J	I	8-11
Purpurine in paste	Co. By	17
Purpurine S in paste		
Purpurine brillante R		3
Dumnwing brillents 4 D	DE	3
Purpurine brillante 4 R	K F	10 107
Pyramine Orange		12—12b
Pyronine B	L. S B	3-8-12
Pyronine GBy.	L, SB	3-8-12
Pyrosine B	Мо	3-17
Pyrosine J	Мо	3
Pyrotine Orange	D	3
Pyrotine R R O		
Rauracienne		:
Red B		
Red C = Soudan III.		
Red Coralline		_
Red Violet R S		3
Red Violet 4 R S		3
Red Violet 5 R extra		3-8-11
Red Violet 5 R S	В	3
Regina Purple	Br S	1
Regina Violet	Br S	1
Regina Violet alcohol soluble		3—8
Regina violet water soluble		3-8
Resorcine Blue		30
Resorcine Brown	Α	•
Resorcine Yellow	E DE	3
December Wielet	A, K F	3—8
Resorcine Violet		
Resorgyl Yellow	K F	12

Rheonine	Name of Dye.	Agent.	Method.
Rhodamine 13			
Rhodamine B extra B, I 3—8—11			
Rhodamine B extra			
Rhodamine 3 B	Phodamine D	D I	
Rhodamine G extra			
Rhodamine 3 G			
Rhodamine 3 G			
Rhodamine 5 G			
Rhodamine 6 G			•
Rhodamine O			
Rhodamine extra B			
Rhodamine B extra B, Klp 3-8-11			
Rhodumine S	Dhalanine extra D	B, Kip	
Rhodindine (see Azo Carmine G paste [B]). By 11 Rhoduline Red G By 11 Rhoduline Red S By 11 Rhoduline Violet By 11 Rhoduline Violet By 11 Roceelline B K, C, D H, G, I, L P, P, P C, R F, S, tM 3 Rock Scarlet Y S Br S 3 Rosazurine B R By 12e Rosazurine B B By 12e Rosazurine B B By 12e Rose Bengale B F 3 Rose Bengale B F 3 Rose Bengale B, C, S 3 Rose Bengale A T A, B 3 Rose Bengale A T A, B 3 Rose Bengale A D M 3 Rose Bengale B M 3 Rose Bengale A C, M Ly 3 Rose Bengale A B 3 Rose Bengale B M 3 Rose Bengale A B 3 Rose Bengale A B 3 Rose Bengale B B 4 Ro	Phodamina C	D. D. J	
Rhoduline Red B	Phodiuding (see Ang Compine Course (D1)	в, ву, г	11
Rhoduline Red G By 11 Rhoduline Red S By 11 Rhoduline Violet By 11 Rocelline B K, C, D H, G, I, L P, P, P C, R F, S, tM 3 Rock Searlet Y S Br S 3 Rock Searlet Y S Br S 3 Rose Searlet B S By 12e Rosazurine B B By 12e Rosazurine B B By 12e Rosazurine B B By 12e Rose Bengale			
Rhoduline Red S			
Rhoduline Violet			
Roceelline B K, C, D H, G, I, L P, P, P C, R F, S, tM 3 Rock Scarlet Y S Br S 3 Rosazine P 3 Rosazurine By 12e Rosazurine B By 12e Rose Bengale BF 3 Rose Bengale BF 3 Rose Bengale BF 3 Rose Bengale A T AB 3 Rose Bengale G M 3 Rose Bengale N C, M Ly 3 Rose de Benzoyle P 12 Rose Diamine B extra Diamine Pink B extra [C] M Ly 3 Rose J B à l'alcool J Rose Magdala D H Roseine B, G, 3 G, R M Rosinduline 2 B bluish K Rosinduline 2 B<	Phoduling Wiglet	By	
Rock Scarlet Y S Br S 3 Rosazine P 3 Rosazurine By 12e Rosazurine B By 12e Rosazurine B B By 12e Rosazurine G By 12e Rose B à l'eau I 3-17 Rose Bengale B F 3 Rose Bengale B, C, S 3 Rose Bengale M, B 3 Rose Bengale A T A, B 3 Rose Bengale G M 3 Rose Bengale N C, M Ly 3 Rose de Benzoyle P 12 Rose Diamine B extra = Diamine Pink B extra [C] M Rose J B à l'aleool J Rose Magdala D H Roseine Br S 711 Roseline B, G, 3 G, R M Rosinduline 2 B bluish K 3 Rosinduline 2 G K 3 Rosolan M 3 Rosophenoline Cl O 12 <td>Pagalling P. F. C. D. H. C. L. T. D. D. D. C.</td> <td>D. D. G. AM</td> <td></td>	Pagalling P. F. C. D. H. C. L. T. D. D. D. C.	D. D. G. AM	
Rosazurine By Rosazurine B By Rosazurine B B By Rosazurine B B By Rosazurine G By Rose B à l'eau I Rose Bengale B F Rose Bengale B, C, S Rose Bengale A T A, B Rose Bengale A T A, B Rose Bengale G M Rose Bengale G M Rose Bengale N C, M Ly Rose Diamine B extra = Diamine Pink B extra [C] Rose J B à l'alcool J Rose Magdala D H Rose Magdala D H Roseine B, G, 3 G, R M Rosinduline 2 B bluish K Rosinduline 2 G K Rosinduline 2 G K Rosolan P Rosolon P <	Parls Constant V.C.	KF, S, IM	
Rosazurine By 12e Rosazurine B By 12e Rosazurine G By 12e Rose B à l'eau I 3—17 Rose Bengale B F 3 Rose Bengale B, C, S 3 Rose Bengale A T A, B 3 Rose Bengale A T A, B 3 Rose Bengale G M 3 Rose Bengale N C, M Ly 3 Rose de Benzoyle P 12 Rose Diamine B extra = Diamine Pink B extra [C] M Rose Magdala D H D Rose Magdala D H N Roseine B, G, 3 G, R M M Rosinduline 2 B bluish K 3 Rosinduline 2 G K 3 Rosolan M 3 Rosolan P 3—7 Rosol Acid Rosophenoline Cl O 12 Rosophenoline M Ly 3 Rouge azoïquet = Azo Red A [C] M Ly 3			
Rosazurine B By 12e Rosazurine B B By 12e Rosazurine G By 12e Rose Ba l'eau I 3—17 Rose Bengale B F 3 Rose Bengale B, C, S 3 Rose Bengale A T A, B 3 Rose Bengale 3 B M 3 Rose Bengale G M 3 Rose Bengale N C, M Ly 3 Rose de Benzoyle P 12 Rose Diamine B extra = Diamine Pink B extra [C] M Ly Rose Magdala D H N Rose Magdala D H Roseine B, G, 3 G, R M Rossinduline 2 B bluish K 3 Rosinduline 2 G K 3 Rosolan M 3 Rosolan P 3—7 Rosol Acid Rosophenoline Cl O 12 Rosophenoline M Ly 3 Rouge azoïquet = Azo Red A [C] M Ly 3			3
Rosazurine B B By 12e Rose B à l'eau I 3—17 Rose Bengale B F 3 Rose Bengale B, C, S 3 Rose Bengale A T A, B 3 Rose Bengale A T A, B 3 Rose Bengale B M 3 Rose Bengale G M 3 Rose Bengale N C, M Ly 3 Rose de Benzoyle P 12 Rose Diamine B extra = Diamine Pink B extra [C] M Rose Magdala D H N Rose Magdala D H Roseine Br S Roseline B, G, 3 G, R M M Rosinduline 2 B bluish K 3 Rosinduline 2 G K 3 Rosolan M M Rosolan P 3—7 Rosol Acid Rosophenoline Cl O 12 Rosophenoline M Ly 3 Rouge I M Ly 3 Rosophenoline M Ly <			40
Rosazurine G By 12e Rose B à l'eau I 3—17 Rose Bengale B F 3 Rose Bengale B F 3 Rose Bengale A T A, B 3 Rose Bengale A T A, B 3 Rose Bengale A T M 3 Rose Bengale A T M 3 Rose Bengale G M 3 Rose Bengale N C, M Ly 3 Rose Bengale N C, M Ly 3 Rose Diamine B extra = Diamine Pink B extra [C] M Ly Rose J B à l'alcool J A Rose Magdala D H B B Rose Magdala D H B B 7—11 Roseline B, G, 3 G, R M M 3 8 Rosinduline 2 B bluish K 3 8 Rosinduline 2 G K 3 8 Rosolan M 3 9 Rosolohenine Cl O 12 Rosophenoline			12e
Rose B à l'eau 3—17 Rose Bengale B F 3 Rose Bengale B, C, S 3 Rose Bengale A T A, B 3 Rose Bengale A T M 3 Rose Bengale B M 3 Rose Bengale G M 2 Rose Bengale N C, M Ly 3 Rose Bengale N C, M Ly 3 Rose Bengale G M 3 Rose Bengale A T M 3 Rose Bengale A M 3 Rose Bengale A M 3 Rose Bengale A B 1 Rose Bengale A B 1	Possagnine C	By	10
Rose Bengale .B F 3 Rose Bengale .B, C, S 3 Rose Bengale A T .A, B 3 Rose Bengale 3 B .M 3 Rose Bengale G .M 3 Rose Bengale N .C, M Ly 3 Rose de Benzoyle .P 12 Rose Diamine B extra = Diamine Pink B extra [C] .M Ly Rose J B à l'alcool .J .J Rose Magdala .D H .D Roseine .Br S .7-11 Roseine B, G, 3 G, R .M Rosinduline 2 B bluish .K 3 Rosinduline 2 G .K 3 Rosolan .M Rosolan .P	Pege D & Page		
Rose Bengale B, C, S 3 Rose Bengale A T A, B 3 Rose Bengale 3 B M 3 Rose Bengale G M 3 Rose Bengale N C, M Ly 3 Rose Ge Benzoyle P 12 Rose Diamine B extra = Diamine Pink B extra [C] M Ly Rose J B à l'alcool J Rose Magdala Rose Magdala D H Roseine Br S Roseline B, G, 3 G, R M M Rosinduline 2 B bluish K 3 Rosinduline 2 G K 3 Rosolan M M Rosolan P 3-7 Rosol Acid P 3-7 Rosophenoline Cl O 12 Rosophenoline M Ly 3 Rouge I M Ly 3 Rouge azoïquet = Azo Red A [C] M Ly 3			
Rose Bengale A T A, B 3 Rose Bengale 3 B M 3 Rose Bengale G M 3 Rose Bengale N C, M Ly 3 Rose Bengale N C, M Ly 3 Rose de Benzoyle P 12 Rose Diamine B extra = Diamine Pink B extra [C] M Ly Rose J B à l'alcool J Rose Magdala D H Roseline B, G, 3 G, R M Rosinduline 2 B bluish K 3 Rosinduline 2 G K 3 Rosinduline 2 G K 3 Rosolan M 3 Rosolan P 3-7 Rosol Acid Rosophenoline Cl O 12 Rosophenoline M Ly 3 Rouge azoïquet = Azo Red A [C] M Ly 3			
Rose Bengale 3 B M 3 Rose Bengale G M 3 Rose Bengale N C, M Ly 3 Rose Bengale N P 12 Rose Bengale N P 12 Rose Bengale N P 12 Rose Diamine B extra = Diamine Pink B extra [C] M Ly Rose J B à l'alcool J A Rose Magdala D H B Roseine S 7-11 Roseline B, G, 3 G, R M M 3-8 Rosinduline 2 B bluish K 3-8 Rosinduline 2 G K 3-8 Rosinduline 2 G K 3-8 Rosolan M M 3-7 Rosolan P 3-7 Rosol Acid Rosophenoline Cl O 12 Rosophenoline Rosophenoline N 3 Rouge I M Ly 3 Rouge azoïquet = Azo Red A [C] M Ly 3			
Rose Bengale G M 3 Rose Bengale N C, M Ly 3 Rose de Benzoyle P 12 Rose Diamine B extra = Diamine Pink B extra [C] M Ly Rose J B à l'aleool J N Rose Magdala D H N Roseïne Br S 7-11 Roseine B, G, 3 G, R M N Rosinduline 2 B bluish K 3 Rosinduline G K 3-8 Rosinduline 2 G K 3 Rosolan M 3-7 Rosol Acid Rosophenine Cl O 12 Rosophenoline Rouge I M Ly 3 Rouge azoïquet = Azo Red A [C] M Ly 3	Pose Dengale 2 D	A, B	
Rose Bengale N C, M Ly 3 Rose de Benzoyle P 12 Rose Diamine B extra = Diamine Pink B extra [C] M Ly Rose J B à l'alcool J Rose Magdala D H Roseïne Br S Roseline B, G, 3 G, R M Rosinduline 2 B bluish K 3 Rosinduline G K 3 Rosinduline 2 G K 3 Rosolan M M Rosolan P 3-7 Rosophenine Cl O 12 Rosophenoline M Ly 3 Rouge I M Ly 3 Rouge azoïquet = Azo Red A [C] M Ly 3	Pose Bengale C	M	
Rose de Benzoyle P 12 Rose Diamine B extra = Diamine Pink B extra [C] M Ly Rose J B à l'alcool J Rose Magdala D H Roseïne Br S Roseline B, G, 3 G, R M Rosinduline 2 B bluish K 3 Rosinduline G K 3 Rosinduline 2 G K 3 Rosolan M M Rosolan P 3-7 Rosophenine Cl O 12 Rosophenoline M Ly 3 Rouge I M Ly 3 Rouge azoïquet = Azo Red A [C] M Ly 3	Pose Dengale W	CMIT	
Rose Diamine B extra [C] M Ly Rose J B à l'aleool J Rose Magdala D H Roseïne Br S 7-11 Roseline B, G, 3 G, R M Rosinduline 2 B bluish K 3 Rosinduline G K 3-8 Rosinduline 2 G K 3 Rosolan M 3-7 Rosolan P 3-7 Rosophenine Cl O 12 Rosophenoline N 12 Rouge I M Ly 3 Rouge azoïquet = Azo Red A [C] M Ly 3			
M Ly Rose J B à l'alcool	Rose Diamine R avtre — Diamine Dink P	orten [C]	12
Rose J B à l'alcool J Rose Magdala D H Roseïne Br S Roseline B, G, 3 G, R M Rosinduline 2 B bluish K 3 Rosinduline G K 3-8 Rosinduline 2 G K 3 Rosolan M 3 Rosolan P 3-7 Rosol Acid P 3-7 Rosophenoline Cl O 12 Rosophenoline M Ly 3 Rouge I M Ly 3 Rouge azoïquet = Azo Red A [C] M Ly 3	Rose Diamine B extra — Diamine I lik B		
Rose Magdala D H Roseïne Br S Roseline B, G, 3 G, R M Rosinduline 2 B bluish K Rosinduline G K Rosinduline 2 G K Rosolan M Rosolan P Rosolan P Rosophenine Cl O Rosophenoline M Rouge I M Ly Rouge azoïquet = Azo Red A [C] M Ly	Rose J. R. à l'algool	M 145	
Roseïne Br S 7—11 Roseline B, G, 3 G, R M M Rosinduline 2 B bluish K 3 Rosinduline G K 3-8 Rosinduline 2 G K 3 Rosolan M P Rosolan P 3-7 Rosol Acid Cl O 12 Rosophenine Cl O 12 Rosophenoline M Ly 3 Rouge I M Ly 3 Rouge azoïquet = Azo Red A [C] M Ly 3			
Roseline B, G, 3 G, R M Rosinduline 2 B bluish K 3 Rosinduline G K 3-8 Rosinduline 2 G K 3 Rosolan M 3 Rosolan P 3-7 Rosol Acid 12 Rosophenine Cl O 12 Rosophenoline , Rouge I M Ly 3 Rouge azoïquet = Azo Red A [C] M Ly 3			7_11
Rosinduline 2 B bluish K 3 Rosinduline G K 3-8 Rosinduline 2 G K 3 Rosolan M M Rosolan P 3-7 Rosol Acid Cl O 12 Rosophenoline Cl O 12 Rosophenoline M Ly 3 Rouge I M Ly 3 Rouge azoïquet = Azo Red A [C] M Ly 3	Roseline R G 3 G R	M	111
Rosinduline G K 3-8 Rosinduline 2 G K 3 Rosolan M M Rosolan P 3-7 Rosol Acid Cl O 12 Rosophenoline Cl O 12 Rosophenoline M Ly 3 Rouge I M Ly 3 Rouge azoïquet = Azo Red A [C] M Ly 3	Rosinduline 2 R bluish	K	2
Rosinduline 2 G K 3 Rosolan M M Rosolan P 3-7 Rosol Acid Cl O 12 Rosophenine Cl O 12 Rosophenoline M Ly 3 Rouge I M Ly 3 Rouge azoïquet = Azo Red A [C] M Ly	Rosinduline G	K	
Rosolan M Rosolan P 3-7 Rosol Acid Cl O 12 Rosophenine Cl O 12 Rosophenoline M Ly 3 Rouge I M Ly 3 Rouge azoïquet = Azo Red A [C] M Ly 3			
Rosolan .P 3-7 Rosol Acid Rosophenine Rosophenoline Rouge I M Ly 3 Rouge azoïquet = Azo Red A [C] M Ly			J
Rosol Acid Rosophenine Rosophenoline Rouge I Rouge azoïquet = Azo Red A [C] M Ly 3			3_7
Rosophenine Cl O 12 Rosophenoline Rouge I M Ly 3 Rouge azoïquet = Azo Red A [C] M Ly			0-1
Rosophenoline , Rouge I M Ly Rouge azoïquet = Azo Red A [C] M Ly			19
Rouge I			12
Rouge azoïquet = Azo Red A [C]	Rouge I	M Lv	3
Rouge B	Rouge azoïquet $=$ Azo Red A [C].	M Ly	U
	Rouge B	M Lv	3—8

· · · · · · · · · · · · · · · · · · ·		
Name of Dye.	Agent.	Method.
Rouge Cerasine	M Lv	
Rouge Chromazone		3
Rouge Congo	R F	3-12
Rouge Congo brillante G	R F	
Rouge Congo brillante R	RF	
Rouge Congo 4 R	RF	12e
Rouge d'acridine B, B B, 3 B	LSR	1-0
Rouge de Lyons A	S S	
Rouge de Naples	р	12b
		120
Rouge Diamine B		
Rouge Diamine 3 B		
Rouge Diamine N 0	M Ly	
Rouge d'oxy-amidodiphényle	Mo	
Rouge foulon $G = Fulling Red G [C]$		
Rouge Français		
Rouge Kyrmezine		
Rouge M	Мо	12
Rouge neutre extra	M Ly	
Rouge Rubis A	M Ly	
Rouge solide Diamine F	M Ly	
Roxamine	D H	
Royal Green crystals	K B	
Rubeosine		
Rubianite		
Rubidine		3
Rubin		7-11
Ruby Ia small crystals		
Rubin S		3-8
Rubramine		11
Rufigallol		4
Russian Green		6a
Russian Leather Red	PSNI	Ott
Russian Leather Red	, 1 3, 1 0	
Saffraniline	C	3-8-11
Saffranine		11
Saffranine A G extra		11
Saffranine A G T extra		11
Saffranine conc		11
Saffranine extra G		11
Saffranine F B extra		
Saffranine F F extra No. 0	By	11
Saffranine G extra, G G S, S		11
Saffranine M N		11
Saffranine P K	C, G, F	11
Saffranine Purple		11
Saffranine S 150		11
Sattranine T		11
Saffranine Scarlet		
Saffran Substitute		
Saffrosine	I. Br S	3—S

Name of Dye.	Agent.	Method,
Salicine Yellow	K	12
Salicine Yellow G, 2 G	K	3
Salicyl Orange		
Salieyl Yellow A	• • • • • • • • •	
Salicyl Yellow B		
Salmon Red	Α	101
Salmon Red	B	12b
Salmon Red	N J	11 12
Sambesi Black B		12
Sambesi Black B R	Д	12
Sambesi Black 3 B	A	12
Sambesi Black D		12
Sambesi Black F	A	12
Sambesi Blue B		12
Sambesi Blue B X	A	12
Sambesi Blue R	<u>A</u>	12
Sambesi Blue R X	A	12
Sambesi Brown	ΑΑ	12
Sambesi Gray B		3
Scarlet 50	n	3
Scarlet 6,503	U	3
Scarlet	T	3
Scarlet Cardinal		3
Scarlet E C		3
Scarlet F R		3
Scarlet F R R		3
Scarlet F R, R R		3
Scarlet for Cotton		3
Scarlet for Silk		3
Scarlet G	Α	3
Scarlet G R Scarlet R Scar	Ry	3
Scarlet R	Kohl	3
Scarlet N R	Kem	3
Scarlet N R R	8.8	3
Seal Brown G		12
Silk Blue	M	7—11
Silk Gray O	M	7
Silk Induline B	K	•
Smaragd Green	By K B	
Solferino	by, 11 15	
Solid Blue	0	3
Solid Blue B alcohol soluble	G	· ·
Solid Blue R alcohol soluble	G	
Solid Blue R water soluble	G	
Solid Blue T water soluble		
Solid Green crystals	C. P.S	3-8-11
Solid Green G	D Н	4
Solid Green J	Mo. P	3-8-11
Solid Green J. J. O		3-8-11
Solid Green O	I	
DOTE OF CHECK IN THE PROPERTY OF THE PROPERTY		

Name of Dye.	. AA	Method.
	Agent.	
Solid Green O in paste	C R, M	6a
Solid Violet	н	4—13a
Solid Yellow N		3
Soluble Blue		811
Soluble Blue B		3
Soluble Blue 8 B		8—11
Soluble Blue 10 B		8-11
Soluble Blue X G		11
Soluble Blue X L		11
Sorbin Red		3—4
Soudan I		
Soudan II		
Soudan III		
Soudan G		
Soudan R		
Soudan Black 1, 2, 3	S S	3
Soudan Black B	S S	3
Soudan Black R, 2 R	S S	3
Soudan Brown	A, Fi	
Soudan Red = Magdala Red.		
Stanley Red	Cl Co	
Standard Blue	A P M	. 12
Standard Brown		12
Standard Cutch Shade	A P M	12
Standard Fast Blue	A P M	. 12
Standard Fast Brown	A P M	12
Standard Fast Red	A P M	12
Standard Fast Tan		12
Standard Fast Yellow		12
Standard Red	A P M	12
Standard Yellow		12
Sterosine Gray	H	
Stilben Red	A	12
Sulfamine Brown A	D	4
Sulfamine Brown B		4
Sulfamine Brown D No. 93		4
Sulfanil Yellow		12
Sulfin Colors (see Cachou de Laval)		15
Sulfon Acid Blue B		3d
Sulfon Acid Blue R		3d
Sulfon Azurine		—11—12e
Sulphin	B	12—12b
Sulpho Black G		3d
Sulpho Black R	Bv	3d
Sulpho Cyanine G	Bv	3d—4
Sulpho Cyanine 3 R	By	3d-4
Sulpho Cyanine 5 R & G R		
Sulpho Green		3
Sulpho Yellow S		
Sun Yellow	By, G. L. S B	12
Sun Gold		

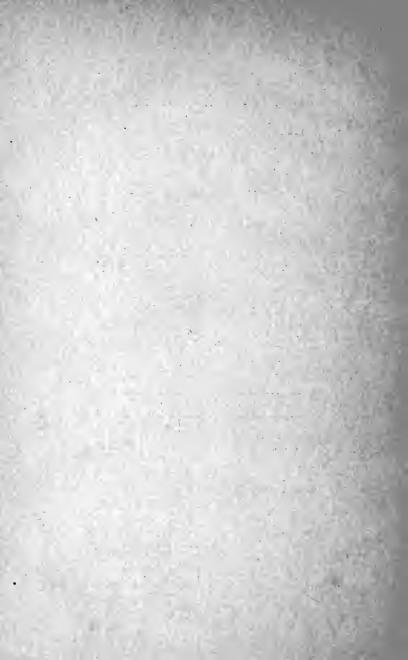
Name of Dye.	Agent.	Method.
Superfine (fine) Magenta crystals	K B	
Superfine Violet	КВ	
Superior Violet	К В	
Superior New Green crystals	K B	
Swiss Black C	S W	12
Swiss Black D	S W	12
Swiss Black B G	S W	12
Swiss Violet R extra	At	3
Tabora Black	A	
Tannin Brown B	C	11
Tannin Heliotrope	C	11
Tannin Indigo	В Н	
Tannin Orange R	C	11
Tartrazine	B. T	3-8
Tennyson Red	H	
Terra cotta F	G	12
Terra cotta R	G	4
Thiamine Yellow	Br S	
Thiazine Brown G	B	12f
Thiazine Brown R	B	12f
Thiazine Red G	В	12f
Thiazine Red R	В	12f
Thiazol Yellow	By	7-11
Thio Brown 2 B	b	
Thio Brown R		12
Thio Carmine R	C	3-8
Thio Carmine R paste	M Lv	3-8
Thio Catechine S		
Thio Catechine No. 1, No. 2, No. 3, No. 4		3
Thio Chromogene	D	12—12b
Thio Cyanosine	Mo	
Thio Flavine T	C, M Ly	11
Thio Flavine S	C. M L7	12b
Thionine		
Thionine Blue G. O extra	M	11
Thio Orange G	D	
Thio Phloxine	Mo	
Thio Phosphine J	L P	
Thio Pyronine		
Thio Ruby	D	3
Thio Vesuvine	Bs	3
Thio Yellow G	Bs	12
Thio Yellow R M	Bs	12
Thio Yellow R		12
Titan Black E D	H	12
Titan Blue B	H	12
Titan Blue B B B	Н	12
Titan Blue S	11	12
Titan Brown O	H	12
Titan Brown R	Н	12

Name of Dye. Agent.	Method.
Tropaeoline Y	3—8
Turkey RedA P M	12
Tumerine	12
Turquois Blue B BBy	
Turquois Blue GBy	
Tyraline	
Tyrian Purple	
Union Black B	1
Union Black B B	1
Union Black SC	12
Union Blue D C	12
Union Blue RA	3—12
Urania BlueD	38
Uranine	38
Ursol DA	
Ursol PA	
Usebe Green	
Cococ dicen	
Vacanceïne Blue	
Vacanceïne Scarlet	
Vat Red Powder	
Vat Red Paste	•
Verde ItalianoL D	12f
Vert acide J J extra concentré M Ly	3—8
Vert brillant = Brilliant Green	3—8—11
Vert de methylaniline = Methyl Green.	3-6-11
Vert de Paris	
Vert Diamant	
Vert Diamine B	
Vert Diamine G = Diamine Green G [C]M Ly	
Vert en pâte	
Vert etincelle	
Vert lumière	
Vert lumière K B	
Vert Naphtol B	
Vert solide cristaux 0 M Ly	
VesuvineB, M	
Vesuvine BB	
Victoria Black BBy	3
Victoria Black GBy	
Victoria Black 5 GBy	
Victoria Blue BB, I	3-8-11
Victoria Blue B S alcohol solubleB, I	3—8—11
Victoria Blue RB, I	3-8-11
Victoria Blue 4 RB, I	3-8-11
Victoria Green 3 BB	3—8—11
Victoria Orange	
Victoria Red	12e
Victoria RubyB K, M	3-8
Victoria Ruby O	3—8

Name of Dye.	Agent	Method.
Victoria Scarlet R	Kchl	3
Victoria Scarlet 3 R		3
Victoria Violet 4 B S		2
Victoria Violet S B S		9
Victoria Yellow		
Victoria Yellow (O Double conc)	M	3
Vidal Black		
Vidal Black S		
Vidaline Blue		12
Vidaline Blue B B		12
Vidaline Blue 5 B		11
Vidaline Blue R		12
Vidaline Blue R R		12
Vidaline Brown B		12
Vidaline Brown G G		12
Vidaline Green		11
Violamine B		3—S
Violamine 3 B		3-8
Violamine G		3-8
Violamine R		3-8
Violaniline		30
Violeine		
Violet A P F		
Violet au Methyl B		3-8-11
Violet au Methyl 6 B		3-8-11
Violet 5 BB		11
Violet 6 BB		11
Violet 7 B		11
Violet Black		3-12
Violet Blue A P		3-1-
Violet C		3-8-11
Violet de methylaniline (see Methyl Violet).	1	3-3-11
Violet de Paris	D	3-8-11
Violet de Faris Violet direct (see Methyl Violet).	1	3311
Violet Diamine N	MITE	12
Violet Formyl S 4 B	MIT	3
Violet neutre extra	MIT	11
Violet neutre solide B		11
Violet non plus ultra	FR	3-8-11
Violet non plus utra	Cz	J
Violet phenilique		3
Violet R		3-8-11
Violet R R		3-8-11
Violet 5 R		3-8-11
Violet 5 R		3-8-11
Violet 3 S O N		3311
Violet 3 S O N		4-13
Violet soluble		3-8-11
Viridine		0-0-11
VIIIUIIIC		

Name of Dye.	Agent.	Method.
Water Blue B		8—11
Water Blue 6 B extra	A D	811
Water Blue B S		811
Water Blue O O		8-11
Water soluble Eosine	M	3—8
White Indephenol = Indephenol White.		,, ,
Welters Bitter = Picrie Aeid.		
Wool Black	A. B	3
Wool Black B		
Wool Black B		3
Wool Black 4 B		3
Wool Black 4 B F	A	3
Wool Black 6 B	A	2-12
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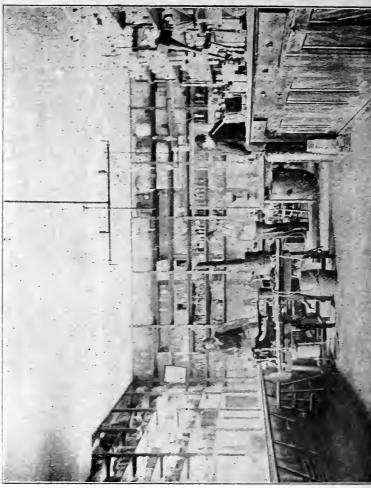
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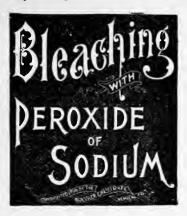
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